

ANGIOGENICALLY EFFECTIVE UNIT DOSE OF FGF-2 AND METHOD OF USE

This application is a continuation-in-part application of U.S. application serial number 09/385,114, filed August 27, 1999, the contents of which are herein incorporated in its entirety.

BACKGROUND OF THE INVENTION

A. Field of the Invention

The present invention is directed to a unit dose composition for inducing cardiac angiogenesis in a human comprising a therapeutically effective amount FGF-2 or an angiogenically active fragment or mutein thereof. The present invention is also directed to a method for administering a single unit dose composition to a human to induce cardiac angiogenesis while minimizing systemic risk to the patient. The present invention is useful because the disclosed unit dose composition, and method for its administration, provide an alternative to angioplasty or surgical intervention for the treatment of coronary artery disease (CAD) and further provide an adjunct for reducing post myocardial infarct (MI) injury in humans.

B. Background of the Invention

The fibroblast growth factors (FGF) are a family of at least eighteen structurally related polypeptides (named FGF-1 to FGF-23) that are characterized by a high degree of affinity for proteoglycans, such as heparin. The various FGF molecules range in size from 15 to at least 32 kD, and exhibit a broad range of biological activities in normal and malignant conditions including nerve cell adhesion and differentiation [Schubert *et al.*, *J. Cell Biol.* 104:635-643 (1987)]; wound healing [U.S. Patent 5,439,818 (Fiddes)]; as mitogens toward many mesodermal and ectodermal cell types, as trophic factors, as differentiation

inducing or inhibiting factors [Clements, *et al.*, *Oncogene* 8:1311-1316 (1993)]; and as an angiogenic factor [Harada, *J. Clin. Invest.*, 94:623-630 (1994)]. Thus, the FGF family is a family of pluripotent growth factors that stimulate to varying extents fibroblasts, smooth muscle cells, epithelial cells and neuronal cells.

5 When FGF is released by normal tissues, such as in fetal development or wound healing, it is subject to temporal and spatial controls. However, many of the members of the FGF family are also oncogenes. Thus, in the absence of temporal and spatial controls, they have the potential to stimulate tumor growth while providing angiogenesis.

10 Coronary artery disease is a progressive condition in humans wherein one or more coronary arteries gradually become occluded through the buildup of plaque (atherosclerosis). The coronary arteries of patients having this disease are often treated by balloon angioplasty or the insertion of stents to prop open the partially occluded arteries. Ultimately, many of these patients are
15 required to undergo coronary artery bypass surgery at great expense and risk. It would be desirable to provide such patients with a medicament that would enhance coronary blood flow so as to reduce the need to undergo bypass surgery.

 An even more critical situation arises in humans when a patient suffers a myocardial infarction, wherein one or more coronary arteries or
20 arterioles becomes completely occluded, such as by a clot. There is an immediate need to regain circulation to the portion of the myocardium served by the occluded artery or arteriole. If the lost coronary circulation is restored within hours of the onset of the infarction, much of the damage to the myocardium that is downstream from the occlusion can be prevented. The clot-
25 dissolving drugs, such as tissue plasminogen activator (tPA), streptokinase, and urokinase, have been proven to be useful in this instance. However, as an adjunct to the clot dissolving drugs, it would also be desirable to obtain collateral circulation to the damaged or occluded myocardium by angiogenesis.

Accordingly, it is an object of the present invention to provide a medicament and a mode of administration that provides human patients with cardiac angiogenesis during coronary artery disease and/or post acute myocardial infarction. More particularly, it is a further object of the present invention to
5 provide a therapeutic dose of an FGF and a mode of administration to humans that provide the desired property of cardiac angiogenesis, while minimizing adverse effects.

Many of the various FGF molecules have been isolated and administered to various animal models of myocardial ischemia with varying and
10 often times opposite results. According to Battler *et al.*, "the canine model of myocardial ischemia has been criticized because of the abundance of naturally occurring collateral circulation, as opposed to the porcine model, which 'excels' in its relative paucity of natural collateral circulation and its resemblance to the human coronary circulation." Battler *et al.*, "*Intracoronary Injection of Basic*
15 *Fibroblast Growth Factor Enhances Angiogenesis in Infarcted Swine Myocardium*," *JACC*, 22(7): 2001-6 (Dec. 1993) at page 2002, col.1. However, Battler *et al.*, who administered bovine bFGF (*i.e.*, FGF-2) to pigs in a myocardial infarct model, considered the varying results that are obtained from one animal species to another, and expressly discloses that the divergent results
20 "thus emphasiz[e] the caution that must be exercised in extrapolating results from different animal models." Battler *et al.*, at page 2005, col.1. Further, Battler points out that "the dosage and mode of administration of bFGF [*i.e.*, bovine FGF-2] may have profound implications for the biologic effect achieved." Battler, *et al.*, at page 2005, col.1. Thus, it is a further object of this invention
25 to discover a dosage and a mode of administration of a fibroblast growth factor that would provide for the safe and efficacious treatment of CAD and/or post MI injury in a human patient. More generally, it is an object of the present invention to provide a pharmaceutical composition and method for inducing angiogenesis in a human heart.

SUMMARY OF THE INVENTION

The Applicants have discovered that administering a single unit dose of about 0.2 µg/kg to about 48 µg/kg of rFGF-2 or an angiogenically active fragment or mutein thereof into one or more coronary vessels (IC) or a peripheral vein (IV) of a human patient in need of coronary angiogenesis, unexpectedly provided the human patient with a rapid and therapeutic coronary angiogenesis that resulted in an unexpectedly large increase (*i.e.*, 96 and 100 seconds of increase in the mean change from baseline for all groups at 2 and 6 months) in the treated patient's exercise tolerance time (ETT) that persisted for an unexpectedly long duration (*i.e.*, 6 months as of this writing). These changes should result in a decreased need for standard revascularization procedures. By the term "coronary angiogenesis," as used herein, is meant the formation of new blood vessels, ranging in size from capillaries to arterioles which act as collaterals in coronary circulation. By way of comparison, angioplasty is considered a therapeutic success if it provides an increase in a patient's ETT of greater than 30 seconds compared to the placebo.

Accordingly, in one aspect, the invention is directed to a unit dose of rFGF-2 comprising a safe and therapeutically effective amount of rFGF-2 or an angiogenically active fragment or mutein thereof. Typically, the safe and therapeutically effective amount comprises about 0.2 µg/kg to about 48 µg/kg of rFGF-2 or an angiogenically active fragment or mutein thereof, based upon ideal body weight. In other embodiments, the safe and therapeutically effective amount of the unit dose comprises 0.2 µg/kg to 2.0 µg/kg, greater than 2.0 µg/kg to less than 24 µg/kg, or 24 µg/kg to 48 µg/kg IC of rFGF-2 or an angiogenically active fragment or mutein thereof. In another embodiment, the safe and therapeutically effective amount of the unit dose comprises 18 µg/kg to 36 µg/kg IV of rFGF-2 or an angiogenically active fragment or mutein thereof. Expressed in absolute terms, the unit dose of the present invention comprises .008 mg to 7.2 mg, more typically 0.3 mg to 3.5 mg, of FGF-2 or an

angiogenically active fragment or mutein thereof. A suitable FGF-2 is the rFGF-2 of SEQ ID NO: 2 or an angiogenically active fragment or mutein thereof.

In another aspect, the present invention is directed to a method of treating a human patient for CAD or to induce coronary angiogenesis therein.

- 5 The method comprises administering into one or more coronary vessels or a peripheral vein of a human patient in need of treatment for coronary artery disease (or in need of angiogenesis) a safe and therapeutically effective amount of a recombinant FGF-2 (rFGF-2) or an angiogenically active fragment or mutein thereof. Typically, a portion of the safe and therapeutically effective amount is
10 administered to each of 2 coronary vessels. The safe and therapeutically effective amount comprises about 0.2 µg/kg to about 48 µg/kg, of rFGF-2 or an angiogenically active fragment or mutein thereof in a pharmaceutically acceptable carrier. In other embodiments, the safe and therapeutically effective amount comprises 0.2 µg/kg to 2 µg/kg, >2 µg/kg to <24 µg/kg, or 24 µg/kg to 48
15 µg/kg of rFGF-2 an angiogenically active fragment or mutein thereof in a pharmaceutically acceptable carrier. In absolute terms, the amount of rFGF-2 or angiogenically active fragment or mutein thereof that is used in the above method comprises .008 mg to 7.2 mg, more typically 0.3 mg to 3.5 mg, of rFGF-2 or an angiogenically active fragment or mutein thereof.

- 20 Because FGF-2 is a glycosoaminoglycan (*e.g.*, heparin) binding protein and the presence of a glycosoaminoglycan optimizes activity and AUC (see Figs. 3 and 4), the IC dosages of RFGF-2 of the present invention typically are administered from 0-30 minutes prior to the administration of a glycosoaminoglycan, such as a heparin. The heparin is administered IC or IV,
25 typically IV. Optionally, the heparin is combined with the unit dose composition.

Because rFGF-2 releases nitric oxide, a potent vasodilator, aggressive fluid management prior to (proactively) and during the infusion is critical to patient's safety. Administration of IV fluids (*e.g.*, 500-1000 mL of normal saline) to establish an estimated wedge pressure of 12 mm Hg prior to

infusion and administration of boluses of IV fluids (e.g., 200 mL normal saline) for decreases of systolic blood pressure (e.g., <90 mm Hg) associated with infusion optimized the safety of administration of rFGF-2 by IC or IV infusion to human patients.

5 Because EDTA is a potent chelator of calcium which is required for normal myocardial contraction and cardiac conduction, minimizing the concentration of EDTA is critical to patient's safety. A concentration of EDTA less than 100 µg/ml in the unit dose composition optimized the safety of administration of rFGF-2 by IC or IV infusion to human patients.

10 Because a sudden bolus of rFGF-2 is associated with profound hypotension in animals, the rate of infusion is critical to patient's safety. Administration at 0.5 to 2 mL per minute, typically 1 mL per minute, optimized the safety of administration of rFGF-2 by IC or IV infusion to human patients.

 The unexpected magnitude and duration of the therapeutic benefit
15 that was provided to human patients in need of coronary angiogenesis by the unit dose composition and method of administration was seen as early as 2 weeks after the single unit dose was administered, and persisted for 6 months after the single unit dose was administered IC or IV, as determined by measuring art-recognized clinical endpoints such as ETT, the "Seattle Angina Questionnaire"
20 (SAQ) and MRI of the target areas of the heart. See, for example, Spertus et al. (1995) *JACC* 25:333-341. In particular, when the ETT of 58 human CAD patients was assessed by treadmill at baseline, and at 1 month, 2 months, and 6 months after administration of a single unit dose of rFGF-2 by IC or IV routes, clinical benefit was observed in some patients in all dosage groups. See Table 1.
25 Increases in exercise capacity appear between 1 and 2 months. The mean ETT increased to greater than 60 seconds at 2 and 6 months with greater benefit being seen in the higher dose group (24-48 µg/kg) than in the mid (6-12 µg/kg) or low (0.33-2.0 µg/kg) dose groups. (See Table 1.) Particularly unexpected and unpredicted by animal models, were the mean increases in ETT in human

patients of 93.4 and 87.5 seconds that were observed at 2 and 6 months, respectively, post-dosing for those patients administered a unit dose of rFGF-2 by IV. Even assuming a placebo effect, the mean change from baseline for the ETT seconds still allowed an unexpectedly favorable comparison of results with angioplasty.

When the quality of life of 48 human CAD patients were assessed by a validated, disease specific questionnaire, the Seattle Angina Questionnaire (SAQ), at baseline (*i.e.*, prior to dosing), and at 2 and 6 months after a single receiving a single unit dose of rFGF-2 of the present invention by IC or IV routes, the mean change from baseline for the 5 scales measured by the SAQ increased in a clinically significant manner for all dosage ranges whether administered IC or IV. (Tables 2-6). In particular, the five scales assessed by the SAQ are exertional capacity, angina stability, angina frequency, treatment satisfaction, and disease perception. Relative to the baseline, the mean score for exertional capacity increased by 10.9 to 20.2 at 2 months; and by 16.5 to 24.1 at 6 months. For angina stability, the mean score increased by 32.1 to 46.2 at 2 months; and by 16.7 to 23.2 at 6 months. For angina frequency, the mean score increased by 20.0 to 32.9 at 2 months; and by 11.4 to 36.7 at 6 months. For treatment satisfaction, the mean score increased by 8.5 to 19.8 at 2 months; and by 6.3 to 19.8 at 6 months. For disease perception, the mean score increased by 20.2 to 27.8 at 2 months; and by 23.8 to 34.0 at 6 months. Generally, a change of 8 points on any scale is considered clinically significant. Thus, the observed changes of 8.5-46.2 are clinically significant for each of the five scales that were assessed. Even assuming a placebo effect whereby a mean change from baseline of 14 points is considered clinically significant, the results still provide for an unexpectedly superior effect at almost all scales that were assessed.

As part of this study, MRI was also performed on 33 human patients diagnosed with CAD to assess the effect of administering a single unit dose of rFGF-2 on their cardiac ejection fraction, regional myocardial function

and perfusion (delayed arrival zone). Specifically, the patients were administered a single unit dose of 0.33 µg/kg to 48 µg/kg IC or 18 µg/kg to 36 µg/kg IV of rFGF-2 of SEQ ID NO: 2. When the 33 human CAD patients were assessed by resting cardiac magnetic resonance imaging (MRI) at baseline (*i.e.*,
5 prior to treatment), and 1, 2 and 6 months after treatment with a single unit dose of rFGF-2 of the invention by IC or IV routes, the patients exhibited a highly statistically significant response to the method of treatment as objectively measured by increased target wall thickening, target wall motion, and target area collateral extent, and by decreased target area delayed arrival extent. (Table 7)
10 By way of summary, at 1, 2 and 6 months, the target wall thickening increased relative to baseline at 4.4%, 6.3% and 7.7%, respectively; the target wall motion increased relative to baseline at 2.7%, 4.4% and 6.4%, respectively; the target area collateral extent increased relative to baseline at 8.3%, 10.9% and 11.2%, respectively; and the target area delayed arrival extent decreased relative
15 to baseline at -10.0%, -8.3% and -10.0%, respectively.

The above data demonstrates the clinical efficacy in humans of the present unit dose composition of rFGF-2 or an angiogenically active fragment thereof when administered IC or IV in accordance with the present invention.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1A is a plot of the mean rFGF-2 plasma concentration versus time profiles for eight different doses of rFGF-2 (SEQ ID NO: 2) administered by IC infusion in humans over a 20 minute period. The eight doses of rFGF-2 presented in Figure 1A are 0.33, 0.65, 2, 6, 12, 24, 36, and 48 $\mu\text{g/kg}$ of lean body mass (LBM).

Figure 1B is a plot of the mean FGF-2 plasma concentration versus time profiles for 2 different doses of rFGF-2 (SEQ ID NO: 2) administered by IV infusion in humans over a 20 minute period. The 2 IV doses of rFGF-2 in Figure 1B are 18 and 36 $\mu\text{g/kg}$. The mean concentration-time profile following IC administration of 36 $\mu\text{g/kg}$ rFGF-2 is included for comparison.

Figure 2 is a plot of mean FGF-2 area under the curve (AUC) in $\text{pg}\cdot\text{min/ml}$ corresponding to Figures 1A and 1B. This plot shows the dose linearity of systemic rFGF-2 exposure following IC or IV infusion. The systemic exposure for the IC route is similar to that observed following IV administration.

Figure 3 is a plot of individual human patient FGF-2 plasma clearance (CL) values as a function of the time of heparin administration in “minutes prior to rFGF-2 infusion” and shows the influence of timing of heparin administration on rFGF-2 plasma clearance (CL).

Figure 4 is a plot individual human patient FGF-2 dose normalized area under curves (AUCs) as a function of the time of heparin administration in “minutes prior to rFGF-2 infusion” and shows the influence of timing of heparin administration on FGF-2 AUC.

Figure 5 summarizes the analysis plan for the Phase II Clinical Trial.

Figure 6 summarizes patient characteristics for the patient population in the Phase II Clinical Trial.

5

Figure 7 shows patient disposition over the course of the Phase II Clinical Trail.

Figure 8 shows the safety variables for the patient population for the
10 Phase II Clinical Trial.

Figure 9 depicts change in exercise time for the placebo, and 3 FGF-2 treatment groups.

15 Figure 10 shows the change in angina frequency score for patients in placebo and the three treatment groups.

Figure 11 shows the change in other domains of the Seattle Angina Questionnaire for patients in placebo and the three treatment groups.

20

Figure 12 shows change in Short Form-36 physical component summary score.

Figure 13 shows change in ETT and angina frequency score stratified by
25 baseline CCS Class 3 or 4.

Figure 14 shows the change in ETT and angina frequency score stratified by baseline AFS less than or equal to 40.

DETAILED DESCRIPTION OF THE INVENTION

The Applicants have discovered that a single dose of rFGF-2 or an angiogenically active fragment or mutein thereof, when administered in a safe and therapeutically effective amount into one or more coronary vessels or into a peripheral vein of a human patient diagnosed with CAD provides the patient with a safe and therapeutically efficacious treatment for the patient's coronary artery disease that lasts at least 4 to 6 months, more typically at least 2 months, before a further treatment is needed. This duration of the effect and the magnitude of the improvements in ETT, SAQ and MRI were unexpected for a single unit dose of medicament.

By the phrase "therapeutically effective amount" or "safe and therapeutically effective amount" as used herein in relation to rFGF-2 is meant an amount of rFGF-2 or an angiogenically active fragment or mutein thereof that when administered in accordance with this invention, is free from major complications that cannot be medically managed, and that provides for objective cardiac improvement in patients having symptoms of CAD despite optimum medical management. Thus, acute hypotension that can be managed by administration of fluids, is considered "safe" for the purpose of this invention. Typically, the safe and therapeutically effective amount of rFGF-2 comprises about 0.2 µg/kg to about 48 µg/kg of rFGF-2 or an angiogenically active fragment or mutein thereof. A suitable FGF-2 for use in the present invention is the rFGF-2 of SEQ ID NO: 2 or an angiogenically active fragment or mutein thereof.

Accordingly, the present invention has multiple aspects. In its first aspect, the present invention is directed to a unit dose composition for inducing angiogenesis in a human patient, the unit dose comprising a therapeutically effective (*i.e.*, an angiogenically effective) amount of rFGF-2 or an angiogenically active fragment or mutein thereof, the amount comprising

about 0.2 µg/kg to about 48 µg/kg of rFGF-2 or an angiogenically active fragment or mutein thereof.

By the term "unit dose composition" as used herein is meant a composition that when administered to a human patient in accordance with the method of the present invention provides a typical human patient in need of angiogenesis with an angiogenic effect of significant efficacy so as not to require retreatment for at least 4-6 months, typically 6 months. The unit dose composition of the present invention is typically provided in combination with one or more pharmaceutically acceptable excipients or carriers. In other embodiments of the unit dose composition, a safe and therapeutically effective amount comprises about 0.2 µg/kg to about 2 µg/kg, about 2 µg/kg to about 24 µg/kg, or about 24 µg/kg to about 48 µg/kg of rFGF-2 or an angiogenically active fragment or mutein thereof.

It is convenient to define the unit dose composition of the present invention in more absolute terms that are not dependent upon the weight of the patient to be treated. When so defined, the unit dose composition comprises from .008 mg to 7.2 mg of rFGF-2 or an angiogenically active fragment or mutein thereof. In this embodiment, the unit dose composition contains a sufficient amount of FGF-2 to accommodate dosing any one of the majority of human CAD patients, ranging from the smallest patient (*e.g.*, 40 kg) at the lowest dosage (about 0.2 µg/kg) through the larger patients (*e.g.*, 150 kg) at the highest dosage (about 48 µg/kg). More typically, the unit dose comprises 0.3 mg to 3.5 mg of rFGF-2 or an angiogenically active fragment or mutein thereof.

The unit dose composition is typically provided in solution or lyophilized form containing the above referenced amount of rFGF-2 and an effective amount of one or more pharmaceutically acceptable buffers, stabilizers and/or other excipients as later described herein.

The active agent in the above described unit dose composition is a recombinant FGF-2 or an angiogenically active fragment or mutein thereof.

Methods for making recombinant FGF-2 are well-known in the art. The recombinant FGF-2 of SEQ ID NO: 2 is made as described in U.S. Patent 5,155,214, entitled "Basic Fibroblast Growth Factor," which issued on October 13, 1992, and which is expressly incorporated herein by reference in its entirety.

5 Moreover, all other references cited herein, whether occurring before or after this sentence, are expressly incorporated herein by reference in their entirety. As disclosed in the '214 patent, a DNA of SEQ ID NO: 1, which encodes a bFGF (hereinafter "FGF-2") of SEQ ID NO: 2, is inserted into a cloning vector, such as pBR322, pMB9, Col E1, pCRI, RP4 or λ -phage, and the cloning vector
10 is used to transform either a eukaryotic or prokaryotic cell, wherein the transformed cell expresses the FGF-2. In one embodiment, the host cell is a yeast cell, such as *Saccharomyces cerevisiae*. The resulting full-length FGF-2 that is expressed has 146 amino acids in accordance with SEQ ID NO: 2. Although the FGF-2 of SEQ ID NO: 2 has four cysteines, *i.e.*, at residue
15 positions 25, 69, 87 and 92, there are no internal disulfide linkages. ['214 at col. 6, lines 59-61.] However, in the event that cross-linking occurred under oxidative conditions, it would likely occur between the residues at positions 25 and 69.

The FGF-2 of SEQ ID NO: 2, which has 146 amino acid residues,
20 differs from naturally occurring human FGF-2 by only 2 amino acid residue. In particular, the amino acids at residue positions 112 and 128 of the FGF-2 of SEQ ID NO: 2 are Ser and Pro, respectively, whereas in human FGF-2, they are Thr and Ser, respectively. In nature, bovine FGF-2, like the corresponding human FGF-2 is initially synthesized *in vivo* as a polypeptide having 155 amino acid
25 residues. Abraham *et al.* "Human Basic Fibroblast Growth Factor: Nucleotide Sequence and Genomic Organization," *EMBO J.*, 5(10):2523-2528 (1986). When the FGF-2 of SEQ ID NO: 2 is compared to the full length 155 residue bovine FGF-2 of Abraham, the FGF-2 of SEQ ID NO: 2 lacks the first nine amino acid residues, Met Ala Ala Gly Ser Ile Thr Thr Leu (SEQ ID NO: 3), at

the N-terminus of the corresponding full length molecule. The recombinant FGF-2 employed in the present compositions and method was purified to pharmaceutical quality (98% or greater purity) using the techniques described in detail in U.S. Pat. 4,956,455, entitled "Bovine Fibroblast Growth Factor" which
5 issued on 09/11/90 and which is incorporated herein by reference in its entirety.

In particular, the first 2 steps employed in the purification of the recombinant FGF-2 of Applicants' unit dose composition are "conventional ion-exchange and reverse phase HPLC purification steps as described previously." [U.S. Pat. 4,956,455, citing to Bolen *et al.*, PNAS USA 81:5364-5368 (1984).] The third
10 step, which the '455 patent refers to as the "key purification step" ['455 at col. 7, lines 5-6], is heparin-SEPHAROSE® affinity chromatography, wherein the strong heparin binding affinity of the FGF-2 is utilized to achieve several thousand-fold purification when eluting at approximately 1.4M and 1.95M NaCl ['455 at col. 9, lines 20-25]. Polypeptide homogeneity was confirmed by
15 reverse-phase high pressure liquid chromatography (RP-HPLC). Buffer exchange was achieved by SEPHADEX® G-25(M) gel filtration chromatography.

In addition to the 146 residue rFGF-2 of SEQ ID NO: 2, the active agent in the unit dose of the present invention also comprises an "angiogenically active fragment" of FGF-2. By the term "angiogenically active
20 fragment" of FGF-2 is meant a fragment of FGF-2 that has about 80% of the 146 residues of SEQ ID NO: 2 and that retains at least 50%, preferably at least 80%, of the angiogenic activity of the FGF-2 of SEQ ID NO: 2.

To be angiogenically active, the FGF-2 fragment should have two cell binding sites and at least one of the two heparin binding sites. The two
25 putative cell binding sites of the analogous human FGF-2 occur at residue positions 36-39 and 77-81 thereof. See Yoshida, *et al.*, "Genomic Sequence of *hst*, a Transforming Gene Encoding a Protein Homologous to Fibroblast Growth Factors and the *int-2*-Encoded Protein," PNAS USA, 84:7305-7309 (Oct. 1987) at Fig. 3. The two putative heparin binding sites of hFGF-2 occur at residue

positions 18-22 and 107-111 thereof. See Yoshida (1987) at Fig. 3. Given the greater than 98% similarity between the amino acid sequences for naturally occurring human FGF-2 (hFGF-2) and rFGF-2 (SEQ ID NO: 2), it is expected that the 2 cell binding sites for rFGF-2 (SEQ ID NO: 2) are also at residue
5 positions 36-39 and 77-81 thereof, and that the 2 heparin binding sites are at residue positions 18-22 and 107-111 thereof. Consistent with the above, it is well known in the art that N-terminal truncations of the FGF-2 of SEQ ID NO: 2 do not eliminate its activity in cows. In particular, the art discloses several naturally occurring and biologically active fragments of the FGF-2 that have N-
10 terminal truncations relative to the FGF-2 of SEQ ID NO: 2. An active and truncated bFGF-2 having residues 12-146 of SEQ ID NO: 2 was found in bovine liver and another active and truncated bFGF-2, having residues 16-146 of SEQ ID NO: 2 was found in the bovine kidney, adrenal glands and testes. [See U.S. Pat. 5,155,214 at col. 6, lines 41-46, citing to Ueno, *et al.*, Biochem and
15 Biophys Res. Comm., 138:580-588 (1986).] Likewise, other fragments of the bFGF-2 of SEQ ID NO: 2 that are known to have FGF activity are FGF-2 (24-120)-OH and FGF-2 (30-110)-NH₂. [U.S. Pat. 5,155,214 at col. 6, lines 48-52.] These latter fragments retain both of the cell binding portions of FGF-2 (SEQ ID NO: 2) and one of the heparin binding segments (residues 107-111).
20 Accordingly, the angiogenically active fragments of FGF-2 typically encompass those terminally truncated fragments of FGF-2 that have at least residues that correspond to residues 30-110 of FGF-2 of SEQ ID NO: 2; more typically, at least residues that correspond to residues 18-146 of FGF-2 of SEQ ID NO: 2.

The unit dose of the present invention also comprises an
25 "angiogenically active . . . mutein" of the rFGF-2 of SEQ ID NO: 2. By the term "angiogenically active . . . mutein" as used herein, is meant an isolated and purified recombinant protein or polypeptide that has 65% sequence identity (homology) to any naturally occurring FGF-2, as determined by the Smith-Waterman homology search algorithm (*Meth. Mol. Biol.* 70:173-187 (1997)) as

implemented in MSPRCH program (Oxford Molecular) using an affine gap search with the following search parameters: gap open penalty of 12, and gap extension penalty of 1, and that retains at least 50%, preferably at least 80%, of the angiogenic activity of the naturally occurring FGF-2 with which it has said at least 65% sequence identity. Preferably, the angiogenically active mutein has at least 75%, more preferably at least 85%, and most preferably, at least 90% sequence identity to the naturally occurring FGF-2. Other well-known and routinely used homology/identity scanning algorithm programs include Pearson and Lipman, *PNAS USA*, **85**:2444-2448 (1988); Lipman and Pearson, *Science*, **222**:1435 (1985); Devereaux *et al.*, *Nuc. Acids Res.*, **12**:387-395 (1984); or the BLASTP, BLASTN or BLASTX algorithms of Altschul, *et al.*, *Mol. Biol.*, **215**:403-410 (1990). Computerized programs using these algorithms are also available and include, but are not limited to: GAP, BESTFIT, BLAST, FASTA and TFASTA, which are commercially available from the Genetics Computing Group (GCG) package, Version 8, Madison WI, USA; and CLUSTAL in the PC/Gene program by Intellegenetics, Mountain View CA. Preferably, the percentage of sequence identity is determined by using the default parameters determined by the program.

The phrase "sequence identity," as used herein, is intended to refer to the percentage of the same amino acids that are found similarly positioned within the mutein sequence when a specified, contiguous segment of the amino acid sequence of the mutein is aligned and compared to the amino acid sequence of the naturally occurring FGF-2.

When considering the percentage of amino acid sequence identity in the mutein, some amino acid residue positions may differ from the reference protein as a result of conservative amino acid substitutions, which do not affect the properties of the protein or protein function. In these instances, the percentage of sequence identity may be adjusted upwards to account for the similarity in conservatively substituted amino acids. Such adjustments are well-

known in the art. See, e.g., Myers and Miller, "Computer Applic. Bio. Sci., 4:11-17 (1988).

To prepare an "angiogenically active mutein" of an angiogenic agent of the present invention, one uses standard techniques for site directed mutagenesis, as known in the art and/or as taught in Gilman, *et al.*, *Gene*, 8:81 (1979) or Roberts, *et al.*, *Nature*, 328:731 (1987). Using one of the site directed mutagenesis techniques, one or more point mutations are introduced into the cDNA sequence of SEQ ID NO: 1 to introduce one or more amino acid substitutions or an internal deletion. Conservative amino acid substitutions are those that preserve the general charge, hydrophobicity/hydrophilicity, and/or steric bulk of the amino acid being substituted. By way of example, substitutions between the following groups are conservative: Gly/Ala, Val/Ile/Leu, Lys/Arg, Asn/Gln, Glu/Asp, Ser/Cys/Thr, and Phe/Trp/Tyr. Significant (up to 35%) variation from the sequence of the naturally occurring angiogenic FGF-2 is permitted as long as the resulting protein or polypeptide retains angiogenic activity within the limits specified above.

Cysteine-depleted muteins are muteins within the scope of the present invention. These muteins are constructed using site directed mutagenesis as described above, or according to the method described in U.S. Pat. 4,959,314 ("the '314 patent"), entitled "Cysteine-Depleted Muteins of Biologically Active Proteins." The '314 patent discloses how to determine biological activity and the effect of the substitution. Cysteine substitution is particularly useful in proteins having 2 or more cysteines that are not involved in disulfide formation. Suitable substitutions include the substitution of serine for one or both of the cysteines at residue positions 87 and 92, which are not involved in disulfide formation. Preferably, substitutions are introduced at the FGF-2 N-terminus, which is not associated with angiogenic activity. However, as discussed above, conservative substitutions are suitable for introduction throughout the molecule.

The unit dose composition of the present invention comprises a safe and an angiogenically effective dose of rFGF-2 or an angiogenically active fragment or mutein thereof, and a pharmaceutically acceptable carrier. Typically, the safe and angiogenically effective dose of the pharmaceutical composition of the present invention is in a form and a size suitable for administration to a human patient and comprises (i) 0.2 $\mu\text{g/kg}$ to 48 $\mu\text{g/kg}$ of rFGF-2 or an angiogenically active fragment or mutein thereof, (ii) and a pharmaceutically acceptable carrier. In other embodiments, the safe and angiogenically effective dose comprises 0.2 $\mu\text{g/kg}$ to 2 $\mu\text{g/kg}$, >2 $\mu\text{g/kg}$ to <24 $\mu\text{g/kg}$ or 24 $\mu\text{g/kg}$ to 48 $\mu\text{g/kg}$ of FGF-2 or an angiogenically active fragment or mutein thereof, and a pharmaceutically acceptable carrier. Expressed in absolute terms for the majority of human CAD patients, the unit dose of the present invention comprises .008 mg to 7.2 mg, more typically 0.3 mg to 3.5 mg, of the FGF-2 or an angiogenically active fragment or mutein thereof.

The second recited component of the unit dose composition of the present invention is a "pharmaceutically acceptable carrier." By the term "pharmaceutically acceptable carrier" as used herein is meant any of the carriers or diluents that are well-known in the art for the stabilization and/or administration of a proteinaceous medicament that does not itself induce the production of antibodies harmful to the patient receiving the composition, and which may be administered without undue toxicity. The choice of the pharmaceutically acceptable carrier and its subsequent processing enables the unit dose composition of the present invention to be provided in either liquid or solid form.

When the unit dose composition is in liquid form, the pharmaceutically acceptable carrier comprises a stable carrier or diluent suitable for intravenous ("IV") or intracoronary ("IC") injection or infusion. Suitable carriers or diluents for injectable or infusible solutions are nontoxic to a human

recipient at the dosages and concentrations employed, and include sterile water, sugar solutions, saline solutions, protein solutions or combinations thereof.

Typically, the pharmaceutically acceptable carrier includes a buffer and one or more stabilizers, reducing agents, anti-oxidants and/or anti-oxidant chelating agents. The use of buffers, stabilizers, reducing agents, anti-oxidants and chelating agents in the preparation of protein based compositions, particularly pharmaceutical compositions, is well-known in the art. See, Wang *et al.*, "Review of Excipients and pHs for Parenteral Products Used in the United States," *J. Parent. Drug Assn.*, **34**(6):452-462 (1980); Wang *et al.*, "Parenteral Formulations of Proteins and Peptides: Stability and Stabilizers," *J. Parent. Sci. and Tech.*, **42**:S4-S26 (Supplement 1988); Lachman, *et al.*, "Antioxidants and Chelating Agents as Stabilizers in Liquid Dosage Forms-Part 1," *Drug and Cosmetic Industry*, **102**(1): 36-38, 40 and 146-148 (1968); Akers, M.J., "Antioxidants in Pharmaceutical Products," *J. Parent. Sci. and Tech.*, **36**(5):222-228 (1988); and Methods in Enzymology, Vol. XXV, Colowick and Kaplan Eds., "Reduction of Disulfide Bonds in Proteins with Dithiothreitol," by Konigsberg, pages 185-188. Suitable buffers include acetate, adipate, benzoate, citrate, lactate, maleate, phosphate, tartarate and the salts of various amino acids. See Wang (1980) at page 455. Suitable stabilizers include carbohydrates such as threlose or glycerol. Suitable reducing agents, which maintain the reduction of reduced cysteines, include dithiothreitol (DTT also known as Cleland's reagent) or dithioerythritol at 0.01% to 0.1% wt/wt; acetylcysteine or cysteine at 0.1% to 0.5% (pH 2-3); and thioglycerol at 0.1% to 0.5% (pH 3.5 to 7.0) and glutathione. See Akers (1988) at pages 225 to 226. Suitable antioxidants include sodium bisulfite, sodium sulfite, sodium metabisulfite, sodium thiosulfate, sodium formaldehyde sulfoxylate, and ascorbic acid. See Akers (1988) at pages 225. Suitable chelating agents, which chelate trace metals to prevent the trace metal catalyzed oxidation of reduced cysteines, include citrate, tartarate, ethylenediaminetetraacetic acid (EDTA) in its disodium, tetrasodium, and

calcium disodium salts, and diethylenetriamine pentaacetic acid (DTPA). *See e.g.*, Wang (1980) at pages 457-458 and 460-461, and Akers (1988) at pages 224-227. Suitable sugars include glycerol, trehalose, glucose, galactose and mannitol, sorbitol. A suitable protein is human serum albumin.

5 In liquid form, a typical unit dose composition of the present invention comprises from about 0.001 mg to 8 mg, more typically 0.03 to 5 mg rFGF-2 or an angiogenically active fragment or mutein thereof, dissolved a pharmaceutically acceptable carrier. A suitable pharmaceutically acceptable carrier comprises 10 mM thioglycerol, 135 mM NaCl, 10 mM sodium citrate,
10 and 1 mM EDTA, pH 5. A suitable diluent or flushing agent for the above-described unit dose composition is any of the above-described carriers. Typically, the diluent is the carrier solution. rFGF-2 or an angiogenically active fragment or mutein thereof is unstable for long periods of time in liquid form. To maximize stability and shelf life of the liquid form, the unit dose composition
15 should be stored frozen at -60°C. When thawed, the solution is stable for 6 months at refrigerated conditions. A typical unit dose would comprise about 1-40 ml, more typically 10-40 ml, of the above-described composition having 0.008-7.2 mg of rFGF-2 or an angiogenically active fragment or mutein dissolved therein. A suitable rFGF-2 for use in the unit dose is the rFGF-2 of
20 SEQ ID NO: 2 or an angiogenically active fragment or mutein thereof.

 In another embodiment, the unit dose composition is provided in lyophilized (freeze-dried) form. In this form, the unit dose of rFGF-2 is capable of being stored at refrigerated temperatures for substantially longer than 6 months without loss of therapeutic effectiveness. Lyophilization is accomplished
25 by the rapid freeze drying (*i.e.*, removing water) under reduced pressure of a plurality of vials, each containing the above described liquid form of the unit dose of the rFGF-2 of the present invention therein. Lyophilizers, which perform the above described lyophilization, are commercially available and readily operable by those skilled in the art. The resulting lyophilized unit dose

composition, in lyophilized cake form, is formulated to contain within the resulting lyophilized cake one or more of the buffers, stabilizers, anti-oxidants, reducing agents, salts and/or sugars described above for the corresponding liquid formulation. A lyophilized unit dose composition containing all such other
5 components need only be reconstituted to a known volume or concentration with sterile aqueous diluent such as sterile water, a sterile sugar solution, or a sterile saline solution. Alternatively, it could be reconstituted with a sterile buffer solution as described above, but lacking a chelating agent, such as EDTA. As a lyophilized cake, the unit dose composition is stable from 6 months to 2 years at
10 refrigerated temperatures. Thus, storage of the unit dose composition in lyophilized form is readily accommodated using conventional refrigeration equipment.

Because the unit dose composition of the present invention is administered via a cardiac catheter or other injection device, which has dead
15 space, it is convenient to formulate the vial containing the unit dose composition so that it contains about 10-50% more of the rFGF-2 or angiogenically active fragment or mutein thereof than is to be administered to the patient. For example, when the unit dose of the rFGF-2 to be administered is 7.2 mg, the vial is optionally formulated to contain up to 50% extra (*e.g.*, a total of about 10.8
20 mg) of rFGF-2 or angiogenically active fragment or mutein thereof. The extra solution is suitable for filling the dead space in the delivery equipment. In an alternative embodiment that does not allow for dead space, the pharmaceutical composition is loaded in the cardiac catheter in front of a pharmaceutically acceptable buffer, diluent or carrier, which is then used to deliver the appropriate
25 amount of the one or more dosages to the one or more sites in the myocardium that are in need of angiogenesis.

As discussed above, the pharmaceutically acceptable carrier for the above described unit dose composition comprises a buffer and one or more stabilizers, reducing agents, anti-oxidants and/or anti-oxidant chelating agents. It

is also within the scope of the present invention that the unit dose composition contain an amount of a glycosoaminoglycan (also known as a "proteoglycan" or a "mucopolysaccharide"), such as heparin, that is effective to bind to the FGF-2 and to the endothelial cell receptors so as to enhance the angiogenic effectiveness of the FGF-2 or angiogenically active fragment or mutein thereof. The amount of heparin that is administered is about 10-80 U per kg of patient weight (U/kg), typically about 40 U/kg. Expressed in absolute terms, the total amount of heparin administered to any one patient does not exceed 5,000 U. Thus, upon reconstitution, the unit dose composition of the present invention would not only contain an angiogenically effective amount of rFGF-2 or an angiogenically active fragment or mutein thereof, it would also contain from about 10-80 U/kg of heparin, typically about 40 U/kg. The typical volume of diluent is from about 1 to 40 ml. While larger volumes of diluent could be used, such larger volumes would typically result in longer administration times. Depending upon the weight of the patient in kg, a single dose comprising from 0.2 µg/kg to 48 µg/kg of the rFGF-2 or an angiogenically active fragment or mutein thereof is withdrawn from the vial as reconstituted product for administration to the patient. Thus, an average 70 kg man that is being dosed at 24 µg/kg would have a sufficient volume of the reconstituted product withdrawn from the vial to receive an IC infusion of (70 kg x 30 µg/kg) 2100 µg (*i.e.*, 2.1 mg).

In its second aspect, the present invention is directed to a method for treating a human patient for CAD or MI, using the above described unit dose composition. In particular, in one embodiment, the present invention is directed to a method for treating a human patient for coronary artery disease, comprising administering a safe and therapeutically effective amount of a recombinant FGF-2 or an angiogenically active fragment or mutein thereof to one or more, typically 2, patent coronary vessels or a peripheral vein of a human patient in need of treatment for coronary artery disease. The human patient in need of treatment for coronary artery disease is typically a human patient with coronary

artery disease who remains symptomatic with angina despite optional medical management. A preferred coronary vessel is a coronary artery, although grafted saphenous veins and grafted internal mammary arteries, as provided by coronary angioplasty, are also suitable. Suitable peripheral veins for administering the
5 unit dose composition include those peripheral veins found throughout the human body that are routinely used by treating physicians and nurses for administration of fluids and medicaments. Examples of such veins include the cephalic, the median cubital, and the basilic of the arm.

When administered as an intracoronary (IC) infusion, the unit dose
10 of rFGF-2 or angiogenic fragment or mutein thereof is typically administered within an hour, more typically over a period of about 20 minutes into one or more (typically, 2) patent coronary vessels. When administered over a twenty minute period, the unit dose composition is typically administered at a rate of 0.5 to 2.0 ml/minute, more typically at about 1 ml/minute. The coronary vessels can
15 be native vessels or grafts, so long as they are not occluded. The volume of the unit dose of rFGF-2 or angiogenic fragment or mutein thereof is typically 10-40 ml; more typically 20 ml. The length of time for infusion of the unit dose is not critical and can be shortened or lengthened depending on the rate and volume of infusion

20 When administered as an intravenous (IV) infusion, the unit dose of rFGF-2 or angiogenic fragment or mutein thereof is administered typically within an hour, more typically over a 20 minute period, into a peripheral vein using a conventional IV setup. When administered over a twenty minute period, the unit dose composition is typically administered at a rate of 1 ml/minute.

25 In the phase I clinical trial of the above described method for treating CAD, a single unit dose composition was administered IC or IV to human patients having CAD who remained symptomatic with angina despite optional medical management. Because the method of the present invention induces angiogenesis, the method of the present invention provides treatment of the

underlying condition in CAD or MI and not merely transitory relief from the symptoms, such as provided by nitrates. Typically, the safe and therapeutically effective amount of the method of the present invention comprises 0.2 µg/kg to 48 µg/kg of rFGF-2 or an angiogenically active fragment or mutein thereof in a pharmaceutically acceptable carrier. In other embodiments, the safe and therapeutically effective amount comprises 0.2 µg/kg to 2 µg/kg, >2 µg/kg to <24 µg/kg, or 24 µg/kg to 48 µg/kg of rFGF-2 or an angiogenically active fragment or mutein thereof in a pharmaceutically acceptable carrier. In absolute terms, the safe and therapeutically effective amount is about .008 mg to about 7.2 mg of rFGF-2 or an angiogenically active fragment or mutein thereof; more typically, 0.3 mg to 3.5 mg of rFGF-2 or an angiogenically active fragment or mutein thereof. A suitable rFGF-2 is the rFGF-2 of SEQ ID NO: 2 or an angiogenically active fragment or mutein thereof.

In another aspect, the present invention is also directed to a method for inducing angiogenesis in a heart of a human patient comprising, administering a single unit dose composition of a recombinant FGF-2 or an angiogenically active fragment or mutein thereof to one or more coronary vessels or to a peripheral vein in a human patient in need of coronary angiogenesis, said unit dose composition comprising from about .008 mg to 7.2 mg of recombinant rFGF-2 or an angiogenically active fragment or mutein thereof in a pharmaceutically acceptable carrier. More typically, the unit dose composition comprises about 0.3-3.5 mg rFGF-2 or an angiogenically active fragment or mutein thereof in a pharmaceutically acceptable carrier. As described above, a single unit dose composition containing a therapeutically effective amount of an rFGF-2 or an angiogenically fragment or mutein thereof is administered to at least one coronary vessel of a human patient in need of angiogenesis, using standard cardiac catheterization techniques already known and used in the art for the intracoronary administration of medicaments, *e.g.*, thrombolytics, streptokinase, or radio-opaque dyes or magnetic particles used to visualize the

coronary arteries. By way of example, a coronary catheter is inserted into an artery (*e.g.*, femoral or subclavian) of the patient in need of treatment and the catheter is pushed forward, with visualization, until it is positioned in the appropriate coronary vessel of the patient to be treated. Using standard
5 precautions for maintaining a clear line, the pharmaceutical composition in solution form is administered by infusing the unit dose substantially continuously over a period of 10 to 30 minutes. Although the pharmaceutical composition of the invention could be administered over a longer period of time, the Applicants perceive no benefit and a potentially increased risk of thrombosis in doing so.

10 Typically, a portion (*e.g.*, one half) of the unit dose is administered in a first coronary vessel. Then, the catheter is repositioned into a second secondary coronary vessel and the remainder of the unit dose is administered with flushing of the catheter. Using the above-described repositioning procedure, portions of the unit dose may be administered to a plurality of coronary vessels until the
15 entire unit dose has been administered. After administration, the catheter is withdrawn using conventional art known protocols. In the phase I clinical trials described herein, therapeutic benefit was reported by patients as early as 2 weeks following the IC rFGF-2 administration of a single unit dose. Clinically significant improvement was demonstrable by objective criterion (ETT and/or
20 SAQ) as early as 30 days following IC or IV administration of a single unit dose of the present invention, and was maintained for 2 months following dosing. In certain patients with progressive CAD disease, it may be necessary or appropriate to administer additional unit doses of rFGF-2 at 2 or 12 month intervals after the initial unit dose, to overcome the progression of the CAD
25 during that interim period. In some patients with very progressive CAD, unit doses of the present invention would be readministered at 4 month intervals. In any instance, the treating physician would be able to determine the time, if any, for readministration based upon routine assessment of the clinical symptoms of the patient.

One of the benefits of the method of the present invention is cardiac angiogenesis. Accordingly, in another aspect, the present invention is directed to a method for inducing angiogenesis in a heart of a human patient, comprising administering into one or more coronary vessels (IC) or into a peripheral vein (IV) of a human patient in need of coronary angiogenesis, a single unit dose composition comprising an angiogenically effective amount of rFGF-2 or an angiogenically active fragment or mutein thereof in a pharmaceutically acceptable carrier. In the above method, the angiogenically effective amount comprises about 0.2 $\mu\text{g/kg}$ to about 48 $\mu\text{g/kg}$ (or in absolute terms about .008 mg to about 7.2 mg) of a recombinant FGF-2 or an angiogenically active fragment or mutein thereof. More typically, the angiogenically effective amount comprises about 0.3 mg to 3.5 mg of a recombinant FGF-2 or an angiogenically active fragment or mutein thereof. A suitable rFGF-2 for use in the above-identified method is the rFGF-2 of SEQ ID NO: 2 or an angiogenically active fragment thereof. In one embodiment of the above method, the unit dose composition is administered IC to patent coronary vessels or IV to a peripheral vein. In another embodiment, the unit dose composition is administered with heparin as described herein.

The above described method for providing coronary angiogenesis is also beneficial in human patients that have undergone a myocardial infarction (MI) in one or more coronary arteries. Accordingly, in another aspect, the present invention is also directed to a method for treating a human patient for an MI comprising, administering into one or more coronary vessels or into a peripheral vein of said human patient, a single unit dose composition comprising a therapeutically effective amount of rFGF-2 or an angiogenically active fragment or mutein thereof. In the above method, the unit dose composition typically comprises about 0.2 $\mu\text{g/kg}$ to about 48 $\mu\text{g/kg}$ (or in absolute terms about .008 mg to about 7.2 mg) of a recombinant FGF-2 or an angiogenically active fragment or mutein thereof in a pharmaceutically acceptable carrier. A

suitable rFGF-2 for use in the above-identified method is the rFGF-2 of SEQ ID NO: 2 or an angiogenically active fragment thereof.

5 In the event of unstable angina or acute myocardial infarction, requiring angioplasty, the same doses of rFGF-2 or angiogenic fragment or mutein thereof that are disclosed herein would also be useful as an adjunct therapy in treating those conditions. Accordingly, in another aspect, the present invention is directed to an improved method for treating a patient for unstable angina or acute myocardial infarction, requiring angioplasty, the method comprising providing angioplasty to the patient in need of treatment; the
10 improvement comprising administering into one or more coronary vessels or into a peripheral vein of said human patient, a single unit dose composition comprising a therapeutically effective amount of rFGF-2 or an angiogenically active fragment or mutein thereof. In the above method, the unit dose composition comprises about 0.2 µg/kg to about 48 µg/kg (or in absolute terms
15 about .008 mg to about 7.2 mg) of a recombinant FGF-2 or an angiogenically active fragment or mutein thereof in a pharmaceutically acceptable carrier. A suitable rFGF-2 for use in the above-identified method is the rFGF-2 of SEQ ID NO: 2 or an angiogenically active fragment thereof.

20 In any of the above-described methods of the present invention, the rFGF-2 or the angiogenically active fragment or mutein thereof is associated with release of nitric oxide, a recognized smooth muscle dilator, which upon administration to the patient causes a sudden drop in the patient's blood pressure. Accordingly, in the methods of the present invention, it is preferable to hydrate the patient with IV fluids prior to administering the unit dose of the present
25 invention. Moreover, for safety and tolerability of the unit dose, aggressive fluid management during and after rFGF-2 administration is also preferred. Finally, it is also within the scope of the above described methods to include the step of administering an effective amount of a glycosoaminoglycan (also known as a "proteoglycan" or a "mucopolysaccharide"), such as heparin from 0-30 minutes

prior to administering the unit dose composition of the present invention. Typically, the effective amount of glycosaminoglycan (such as heparin) that is administered is about 10-80 U/kg, more typically, about 40 U/kg. However, the total amount of heparin administered to any one patient immediately prior to
5 dosing generally does not exceed 5,000 U.

Because EDTA is a potent chelator of calcium, which is required for normal myocardial contraction and cardiac conduction, minimizing the concentration of EDTA is critical to patient's safety. A concentration of EDTA less than 100 µg/ml optimized the safety of administration of rFGF-2 by IC or
10 IV infusion to human patients.

Because a sudden bolus of rFGF-2 is associated with profound hypotension in animals, the rate of infusion is critical to patient's safety. Administration at 0.5 to 2 ml per minute, typically 1 ml per minute, optimized the safety of administration of rFGF-2 by IC or IV infusion to human patients.

15 A Phase I clinical trial directed to treating human patients for CAD by administering a single unit dose composition of the present invention was conducted and is described in Examples 1-3 herein. In that trial, 66 human patients diagnosed with CAD, who satisfied the criteria of Example 2 herein, received a single unit dose of rFGF-2 in accordance with the method of the
20 present invention. Specifically, 52 human patients were administered a unit dose of 0.33 µg/kg to 48 µg/kg of rFGF-2 by IC infusion over about a 20 minute period. Fourteen human patients were administered a unit dose of either 18 µg/kg or 36 µg/kg of rFGF-2 by IV infusion over about a 20 minute period. The 66 treated patients were then assessed relative to baseline (*i.e.*, prior to
25 treatment with the single unit dose), and again at 1 month, 2 months and 6 months after treatment with the single unit dose, using three sets of art-recognized assessment criteria: 1) changes in their exercise tolerance time (ETT); 2) the Seattle Angina Questionnaire, which provides an assessment based upon a

mixed combination of objective and subjective criteria; and 3) the measurement of physical changes in the heart as assessed by MRI.

For ETT of the 66 patients of the Phase I clinical trial of Examples 1-3 was measured at baseline, and at 1 month, 2 months and 6 months after dosing (with a single unit dose composition of the invention) using a Bruce treadmill protocol. Subjects were excluded from the analysis if the treadmill protocol was not the same as used at baseline. Therefore, the number of subjects varied over time. In addition, any patients who had emergency revascularization were excluded from the analysis. A dose was considered effective if the mean change in ETT from baseline increased by greater than 60 seconds. The results of the ETT assessment are provided in Table 1.

Table 1. Exercise Tolerance Time (ETT) - Change from Baseline

<i>FGF-2 Dose Group</i>	<i>Change from Baseline at One Month</i>	<i>Change from Baseline at Two Months</i>	<i>Change from Baseline at Six Months</i>
0.33 to 2.0 µg/kg IC (N = 16) (low)	N = 8 45.1 sec (-105 to 180)	N = 6 130.0 sec* (19 to 240)	N = 5 60.8 sec (-45 to 210)
6.0 and 12 µg/kg IC (N = 8) (mid)	N = 2 -24.0 sec (-48 to 0)	N = 4 -2.5 sec (-90 to 120)	N = 2 6.5 sec (-0 to 13)
24.0 to 48.0 µg/kg IC (N = 28) (high)	N = 18 51.9 sec (-188 to 399)	N = 21 107.9 sec* (-30 to 385)	N = 16 133.1 sec* (-195 to 386)
18.0 & 36.0 µg/kg IV (N = 14)	N = 12 45.1 sec (-75 to 237)	N = 12 93.4 sec* (0 to 285)	N = 12 87.5 sec* (-60 to 285)
ALL GROUPS (N = 66)	N = 40 45.0 sec	N = 43 96.0 sec	N = 35 100.0 sec

15 N = number of subjects; mean; (range in seconds); * = p < 0.05

Referring to Table 1, the mean change from baseline at one month was less than 60 seconds for all dose groups. However, the percentage of patients stopping their treadmill test because of angina decreased in all groups over time. At 2

months and 6 months after dosing, the mean changes from baseline were greater in the high dose IC and IV groups of patients than in the low and mid dose IC groups. The persistence of increased ETT at 6 months (133.1 sec and 87.5 sec) in the high dose IC (24-48 :g/kg) and IV (18 & 38 :g/kg) groups, respectively, was unexpected. The greatest mean increases in ETT of 107.9 and 133.1 seconds at 2 and 6 months, respectively, occurred in the high dose (24-48 :g/kg) IC group. The IV group exhibited significant mean increases in ETT of 93.4 seconds and 87.5 seconds, at 2 months and 6 months respectively, which was not predicted by the rat and pig animal models used herein. Overall, the persistence of the effect (increase in ETT) at 2 months and its magnitude for both the IC and IV groups was wholly unexpected.

The 66 human patients of the Phase I clinical trial described in Examples 1-3 herein were also evaluated using the Seattle Angina Questionnaire (SAQ). The SAQ is a validated, disease-specific, quality of life instrument which assesses the following five scales: 1) "exertional capacity" = limitation of physical activity; 2) "disease perception" = worry about MI; 3) "treatment satisfaction"; 4) "angina frequency" = number of episodes and sublingual nitroglycerin usage; and 5) "angina stability" = number of episodes with most strenuous physical activity. The possible range of scores for each of the five scales is 0 to 100 with the higher scores indicating a better quality of life. Typically, a mean change of 8 points or more between the mean baseline scores (*i.e.*, before treatment) and the post-treatment scores is recognized as being "clinically significant." However, in the present analysis, a dose was considered "effective" if the mean change in score from baseline increased by greater than 14 points. The reason that 14 was chosen (instead of 8) was to allow for the improvement that was seen in the placebo group at 2 months in a clinical trial of another growth factor--VEGF.

In performing the SAQ evaluation, the patients were categorized according to the same dosage groups that were evaluated for ETT, *i.e.*, 0.33 -

2.0 µg/kg IC (low) 6.0 - 12.0 µg/kg IC (mid); 24 - 48 µg/kg IC (high); and 18 and 36 µg/kg IV. The questionnaire was administered to subjects in each dosage group at baseline (prior to dosing), and at 2 months and 6 months after being administered a single unit dose composition of rFGF-2 in accordance with the method of the present invention.

The first SAQ scale is "exertional capacity." The data on exertional capacity is summarized in Table 2 herein. As reflected in Table 2,

Table 2. Exertional Capacity (EC) - Change from Baseline

<i>FGF-2 Dose Group</i>	<i>Change from Baseline at Two Months</i>	<i>Change from Baseline at Six Months</i>
0.33 to 2.0 µg/kg IC (N = 16)	N = 14 15.0* (-25 to 53)	N = 7 23.2* (0 to 53)
6.0 and 12 µg/kg IC (N = 8)	N = 7 20.2* (-14 to 44)	N = 6 24.1 (-11 to 69)
24.0 to 48.0 µg/kg IC (N = 28)	N = 26 14.6* (-33 to 75)	N = 23 22.9* (-14 to 75)
18.0 and 36.0 µg/kg IV (N = 14)	N = 12 10.9 (-8 to 67)	N = 14 16.5* (-19 to 63)

N = number of subjects; mean (range); * = p < 0.05

the change from baseline in mean score increased at 2 and 6 months for each of the three IC dosage groups and at 6 months for all dosage groups (IC and IV). All scores at all dosage levels increased with time in going from 2 months to 6 months with the best increases (23.2, 24.1, 22.9 and 16.5) relative to baseline being seen at 6 months post-dosing.

The second SAQ scale to be evaluated was "angina stability." The data summarizing the angina stability is presented in Table 3 herein.

Table 3. Angina Stability (AS) – Change from Baseline

<i>FGF-2 Dose Group</i>	<i>Change from Baseline at Two Months</i>	<i>Change from Baseline at Six Months</i>
0.33 to 2.0 µg/kg IC (N = 16)	N = 13 46.2* (0 to 100)	N = 7 21.4* (0 to 50)
6.0 and 12 µg/kg IC (N = 8)	N = 7 32.1* (0 to 50)	N = 6 16.7 (-25 to 50)
24.0 to 48.0 µg/kg IC (N = 28)	N = 27 34.3* (-25 to 75)	N = 24 17.7* (-25 to 75)
18.0 and 36.0 µg/kg IV (N = 14)	N = 12 39.6* (0 to 100)	N = 14 23.2* (0 to 75)

N = number of subjects; mean (range); * = p < 0.05

According to Table 3, the change in score for angina stability increased relative to baseline at both 2 and 6 months for each group. The improvements in angina stability seen at 2 months after dosing (46.2, 32.1, 34.3 and 39.6) were significantly greater than those scores seen at 6 months (21.4, 16.7, 17.7 and 23.2). However, the scores found at both 2 months and 6 months after dosing showed that all dosages were found to be effective (>14) in increasing angina stability. Moreover, the magnitude of the increases and their duration for 6 months were unexpected.

The third SAQ scale to be evaluated was “angina frequency.” The data summarizing the angina frequency is presented in Table 4 herein.

Table 4. Angina Frequency (AF) – Change from Baseline

<i>FGF-2 Dose Group</i>	<i>Change from Baseline at Two Months</i>	<i>Change from Baseline at Six Months</i>
0.33 to 2.0 µg/kg IC (N = 16) (low)	N = 14 27.9* (-10 to 80)	N = 7 12.9 (-40 to 50)
6.0 and 12 µg/kg IC (N = 8) (mid)	N = 7 32.9* (0 to 80)	N = 6 36.7 (-10 to 90)
24.0 to 48.0 µg/kg IC (N = 28) (high)	N = 27 28.9* (-40 to 80)	N = 24 25.8* (-30 to 80)
18.0 and 36.0 µg/kg IV (N = 14)	N = 12 20.0* (0 to 90)	N = 14 11.4 (-30 to 60)
ALL GROUPS (N = 66)	N = 60 27.3	N = 51 21.4

N = number of subjects; mean (range); * = p < 0.05

According to Table 4, the mean patient scores (27.9, 32.9, 28.9 and 20.0) for angina frequency increased at 2 months (relative to baseline) by an effective amount (>14) for all dosage groups and for all modes of administration (IC or IV). The mean patient scores continued to increase at 6 months only for the mid dose (6.0 - 12.0 :g/kg) group, suggesting a peak effect at 2 months post-dosing. However, for the mid dose (6.0 - 12.0 :g/kg) and high dose (24.0 - 48.0 :g/kg) groups, the changes at 2 months and 6 months were similar, suggesting a persistent effect at 6 months on angina frequency.

The fourth SAQ scale to be evaluated was “treatment satisfaction.” The data summarizing the angina frequency is presented in Table 5 herein.

Table 5. Treatment Satisfaction (TS) – Change from Baseline

<i>FGF-2 Dose Group</i>	<i>Change from Baseline at Two Months</i>	<i>Change from Baseline at Six Months</i>
0.33 to 2.0 µg/kg IC (N = 16) (low)	N = 14 8.5* (-19 to 31)	N = 7 6.3 (-25 to 25)
6.0 and 12 µg/kg IC (N = 8) (mid)	N = 7 17.9 (-13 to 44)	N = 6 19.8 (0 to 63)
24.0 to 48.0 µg/kg IC (N = 28) (high)	N = 27 18.8* (-38 to 69)	N = 24 13.0 (-75 to 63)
18.0 and 36.0 µg/kg IV (N = 14)	N = 12 19.8* (-13 to 63)	N = 14 13.4* (-19 to 56)

N = number of subjects; mean (range); * = p < 0.05

According to Table 5, the score for treatment satisfaction increased by an effective amount at 2 months for the mid and high dose IC groups as well as the IV group. At 2 months post-dosing, only the score for the mid dose group IC had a score that was greater than 14, suggesting a peak effect for treatment satisfaction at 2 months.

The fifth SAQ scale to be evaluated was “disease perception.” The data summarizing the disease perception is presented in Table 6 herein. According to Table 6, the scores for disease perception increased from baseline to scores of 20.2-29.2 at 2 months and 23.8-34.0 at 6 months. These scores showed that administering a single unit dose composition in accordance with the method of the present invention was considered to be as effective (or more effective) at 6 months as at 2 months. These scores suggest a persistence of the effectiveness of the method of the present invention on disease perception out to 6 months following administration of a single unit dose composition.

Table 6. Disease Perception (DP) – Change from Baseline

<i>Dose Group</i>	<i>Change from Baseline at Two Months</i>	<i>Change from Baseline at Six Months</i>
0.33 to 2.0 µg/kg IC (N = 16) (low)	N = 14 29.2* (-8 to 58)	N = 7 26.2* (0 to 42)
6.0 and 12 µg/kg IC (N = 8) (mid)	N = 7 20.2* (-8 to 50)	N = 6 30.6* (0 to 58)
24.0 to 48.0 µg/kg IC (N = 28) (high)	N = 27 27.8* (-33 to 92)	N = 24 34.0* (-33 to 100)
18.0 and 36.0 µg/kg IV (N = 14)	N = 12 22.9* (-8 to 92)	N = 14 23.8* (-8 to 75)

N = number of subjects; mean (range); * = $p < 0.05$

- Up to 60 of the human patients of the Phase I clinical trial described in Examples 1-3 herein were also evaluated using resting magnetic resonance imaging (MRI) scans of their heart. The resting MRI scans were performed on the patients at baseline, and at 1 month, 2 months and 6 months after dosing with a single unit dose composition of the present invention. The doses were considered “effective” based upon statistical significance ($p < 0.05$).
- The objective criteria assessed by the resting MRI scans are the following: (1) ejection fraction; (2) myocardial infarct extent (%); (3) normal wall thickening (4) normal wall motion (%); (5) target wall thickening (%); (6) target wall motion (%); (7) target wall area collateral extent (%); and (8) target area delayed arrival extent (%).
- Based upon the resting MRI, no change in “ejection fraction” was observed at one month for any one group. The mean change from baseline for all groups ($n = 33$) at 1 month was an increase of 2.0% ($p = 0.042$). At 2 months, the mean change from baseline for the low dose IC group ($n=13$) was an increase of 8.1% ($p=0.007$); and for all groups ($n=54$), the mean change from baseline was an increase of 3.8% ($p=0.001$). At 6 months, the mean

change from baseline for the high dose IC group (n=19) was 5.3% (p=0.023); for the IV group (n=3) was 11.1% (p=0.087); and for all groups (n=33) was 5.7% (p=0.001).

Based upon the resting MRI, there was no statistically significant change in the “myocardial infarct extent” (%) for any group, or for all groups in combination at 1 month, 2 months or 6 months post-dosing. When the normal wall motion (%) and normal wall thickening were assessed, there was no statistically significant change from baseline at 1 month, 2 months or 6 months for any one group. However, there was a statistically significant change from baseline in target wall motion for all groups at one (n=60), 2 (n=54) and 2 (n=33) months, which was reflected as a mean increase from baseline of 2.7% (p=0.015), 4.4% (p=<0.001) and 6.4% (p<0.001), respectively. There was also a statistically significant change from baseline in target wall thickening for all groups at one (n=60), 2 (n=54) and 2 (n=33) months, which was reflected as a mean increase from baseline of 4.4% (p=0.015), 6.3% (p=<0.001) and 7.7% (p<0.001), respectively.

The next criteria assessed by MRI was “target area collateral extent” (%). The mean increase from baseline in target area collateral extent for all groups was highly statistically significant at one month (n=31), 2 months (n=27) and 2 months (n=16), wherein the increases were 8.3% (p<0.001), 10.9% (p<0.001) and 11.2% (p<0.001), respectively. The greatest collateral extent increases were observed for the low and mid IC doses, *i.e.*, at one month (10.4% and 18.3%, respectively), 2 months (14.7% and 18.0%, respectively) and 2 months (16.0% and no value for mid dose, respectively), which was wholly unexpected. At one month, 2 months and 2 months post-dosing, the corresponding % increases in target area collateral extent that were observed for the IC high dose group were 6.3%, 8.0% and 9.0%, respectively.

The final criteria assessed by MRI was “target area delayed arrival extent” (%). The mean decrease from baseline in target area delayed arrival

extent for all groups was highly statistically significant at 1 month (n=60), 2 months (n=54) and 6 months (n=34), wherein the decreases were -5.8% (p<0.001), -8.3% (p<0.001) and -10.0% (p<0.001), respectively. The greatest target area delayed extent decreases were observed for the low dose IC group, which was also highly unexpected.

Thus, providing CAD patients with a single IC or IV infusion of rFGF-2 in accordance with the present invention provided the patients with a statistically significant physical improvement as objectively measured by MRI and other conventional criteria.

10

Pharmacokinetics and Metabolism

The molecular structure of FGF-2 contains a positively charged tail that is known to bind to proteoglycan chains (heparin and heparin-like structures) on cell surfaces and on the endothelial wall of the vasculature. See Moscatelli, *et al.*, "Interaction of Basic Fibroblast Growth Factor with Extracellular Matrix and Receptors," Ann. NY Acad. Sci., 638:177-181 (1981).

The kidneys and liver are the major organs for the elimination of rFGF-2. In particular, the kidneys have a protein cutoff of about 60 kD and thus retain serum albumin (MW 60 kD). However, FGF-2 (146 residues) has a molecular weight of about 16.5 kD. Accordingly, renal excretion is to be expected. In a radiolabelled biodistribution study of commercially available bovine FGF-2 (bFGF-2), both the liver and the kidney were shown to contain high counts of the radiolabelled bFGF-2 at 1 hour after IV or IC injection. In a published study, wherein another recombinant iodinated form of bFGF-2 was given to rats, the liver was identified as the major organ of elimination. Whalen *et al.*, "The Fate of Intravenously Administered bFGF and the Effect of Heparin," Growth Factors, 1:157-164 (1989). It is also known that FGF-2 binds in the general circulation to α_2 -macroglobulin and that this complex is internalized by receptors on the Kupffer cells. Whalen *et al.* (1989) and

LaMarre *et al.*, "Cytokine Binding and Clearance Properties of Proteinase-Activated Alpha-2-Macroglobulins," *Lab. Invest.*, **65**:3-14 (1991). Labelled FGF-2 fragments were not found in the plasma, but they were found in the urine and corresponded in size to intracellular breakdown products.

5 In preclinical testing, we determined the pharmacokinetics of rFGF-2 (SEQ ID NO: 2) after intravenous (IV) and intracoronary (IC) administration in domestic Yorkshire pigs, and after IV administration dosing in Sprague Dawley ("SD") rats. The pig models demonstrated linear pharmacokinetics (0.65 µg/kg - 20 µg/kg) IC and IV. The terminal half-life of
10 the FGF-2 in the pig model was 3-4 hours. The rat models demonstrated linear pharmacokinetics over the range of 30-300 µg/kg IV. The terminal half-life of the FGF-2 in the rat model was 1 hour. Both species showed plasma concentration suggesting a two-compartment model.

 Likewise, in humans, the FGF-2 plasma concentrations after IV and/or IC
15 infusion followed a biexponential curve with an initial steep slope and considerable decrease over several log scales (the distribution phase) during the first hour, followed by a more moderate decline (the elimination phase). Fig. 1A provides a plasma concentration versus time curve showing these phases in humans after IC administration of rFGF-2 of SEQ ID NO: 2 as a function of
20 each of the following eight doses: 0.33 µg/kg, 0.65 µg/kg, 2 µg/kg, 6 µg/kg, 12 µg/kg, 24 µg/kg, 36 µg/kg, and 48 µg/kg of lean body mass (LBM). Figure 1A shows the plasma dose linearity for the eight doses of rFGF-2 that were administered by IC infusion over a twenty minute period. Figure 1A also shows a biphasic plasma level decline, *i.e.*, a fast distribution phase during the first
25 hour, followed by an elimination phase with an estimated $T_{1/2}$ of 5-7 hours. The plasma concentrations of FGF-2 of SEQ ID NO: 2 were determined by a commercially available ELISA (R&D Systems, Minneapolis MN) that was marketed for analysis of human FGF-2. The ELISA assay showed 100% cross-reactivity with the rFGF-2 of SEQ ID NO: 2. Other members of the FGF

family, as well as many other cytokines, were not detected by this assay. Further, heparin does not interfere with the assay.

Figure 1B is a plot of the mean FGF-2 plasma concentration as a function of time for 18 $\mu\text{g/kg}$ and 36 $\mu\text{g/kg}$ rFGF-2 administered IV, as compared to 36 $\mu\text{g/kg}$ rFGF-2 administered IC. The plasma concentration versus time profiles in Figure 1B for the 36 $\mu\text{g/kg}$ doses by the IV and IC routes are superimposable. However, a first-pass effect with the IC route is not eliminated.

Figure 2 is a plot of mean FGF-2 area under the curve (AUC) in $\text{pg}\cdot\text{min/ml}$ corresponding to Figures 1A and 1B. Figure 2 shows the dose linearity of systemic rFGF-2 exposure (AUC) following IC or IV infusion. In particular, Figure 2 shows that the systemic exposure to rFGF-2 following administration by the IC and IV routes is substantially similar.

Figure 3 is a plot of individual human patient plasma clearance (CL) values (ml/min/kg) as a function of the time of heparin administration in “minutes prior to rFGF-2 infusion.” Figure 3 shows the influence of timing of heparin administration on FGF-2 plasma clearance (CL). Although Figure 3 shows that administering heparin up to 100 minutes prior to rFGF-2 decreases FGF-2 clearance, the preferred time for administering heparin is from 0-30 minutes prior the rFGF-2 administration, wherein the effect of the heparin on decreasing FGF-2 clearance is greatest.

Figure 4 is a plot individual human patient rFGF-2 dose normalized area under curves (AUCs) as a function of the time of heparin administration in “minutes prior to rFGF-2 infusion” and shows the influence of timing of heparin administration on rFGF-2 AUC. Figure 4 shows that the greatest AUC/dose was achieved when an effective amount of a glycosaminoglycan, such as heparin, was preadministered within 30 minutes or less of IC rFGF-2 infusion, more preferably within 20 minutes or less of IC or IV rFGF-2 infusion. Typically, an effective amount of a glycosaminoglycan is 10-80 U/kg heparin.

The mean pharmacokinetic parameters for rFGF-2 in humans as a function of dosage and mode of administration are summarized in Table 8 herein. Referring to Table 8, the $T_{1/2}$ for FGF-2 in humans was determined to range from 2.2 ± 3.7 hours at low dose ($0.33\text{-}2.0 \mu\text{g/kg}$) IC to 7.0 ± 3.5 hours at a dose of $18\text{-}36 \mu\text{g/kg}$ IV; given the limitations of the assay, the terminal half-life is estimated at 5-7 hours for all groups. The clearances of FGF-2 ranged from 13.2 to 18.2 L/hour/70kg man. Finally, the steady state volume (V_{ss}) was determined to range from 11.3 ± 10.4 L/70kg man to 16.8 ± 10.7 L/70kg man.

Table 8. Mean rFGF-2 PK Parameters in Humans

<i>FGF-2 Dose $\mu\text{g/kg}$</i>	<i>N</i>	<i>Route</i>	<i>CL (L/hr/70kg)</i>	<i>t_{1/2} (h)</i>	<i>V_{ss} (L/70kg)</i>
0.3 - 2	16	IC	18.2 ± 13.4	2.2 ± 3.7	11.3 ± 10.4
6 - 12	8	IC	13.2 ± 7.3	3.1 ± 2.5	12.1 ± 4.9
24 - 48	28	IC	14.7 ± 8.3	6.3 ± 1.8	16.8 ± 10.7
18 - 36	14	IV	13.9 ± 7.9	7.0 ± 3.5	16.4 ± 8.6

10

Although the binding of FGF-2 to heparin-like structures is strong (dissociation constant $\sim 2 \times 10^{-9}$ M), the binding of FGF-2 to a specific tyrosine kinase receptor is approximately 2 orders of magnitude higher (dissociation constant $\sim 2 \times 10^{-11}$ M). Moscatelli *et al.*, (1991). Thus, without being bound to any theory, the complexation of rFGF-2 with a glycosoaminoglycan, such as a heparin, might increase signal transduction and mitogenesis, and/or protect the rFGF-2 from enzymatic degradation.

Further aspects of the invention for the administration of FGF-2 as disclosed herein include, but are not limited to: to improve exercise capacity as measured by ETT treadmill analysis; to improve exercise capacity as measured by exertional capacity domain of the SAQ; to reduce the frequency of angina; and to increase perfusion as measured by MRI. In particular exercise capacity was significantly improved in patients with congestive heart failure.

20

The following examples are offered by way of illustration and not by way of limitation.

1. If the defendant is a minor, the defendant is not liable for the tortious act.

EXAMPLE 1

“Unit Dose of rFGF-2 Employed in the Phase I Clinical Trial”

5 The rFGF-2 of SEQ ID NO: 2 was formulated as a unit dose and pharmaceutical composition and administered to rats, pigs and ultimately to humans in the Phase I clinical trial referenced herein. The various formulations are described below.

10 **The rFGF-2 Unit Dose** was provided as a liquid in 3 cc type I glass vials with a laminated gray butyl rubber stopper and red flip-off overseal. The rFGF-2 unit dose contained 1.2 ml of 0.3 mg/ml rFGF-2 of SEQ ID NO: 2 in 10 mM sodium citrate, 10 mM monothioglycerol, 1 mM disodium dihydrate EDTA (molecular weight 372.2), 135 mM sodium chloride, pH 5.0. Thus, in absolute terms, each
15 vial (and unit dose) contained 0.36 mg rFGF-2. The vials containing the unit dose in liquid form were stored at 2° to 8° C.

The rFGF diluent was supplied in 5 cc type I glass vials with a laminated gray butyl rubber stopper and red flip-off overseal. The rFGF-2 diluent contains 10
20 mM sodium citrate, 10 mM monothioglycerol, 135 mM sodium chloride, pH 5.0. Each vial contained 5.2 ml of rFGF-2 diluent solution that was stored at 2° to 8°C.

The rFGF-2 Pharmaceutical Composition that was infused was prepared by
25 diluting the rFGF-2 unit dose with the rFGF diluent such that the infusion volume is 10-40 ml. In order to keep the EDTA concentration below a preset limit of 100 µg/ml, the total infusion volume was increased to a maximum of 40 ml when proportionately higher absolute amounts of FGF-2 were administered to patients with higher body weights.

30

EXAMPLE 2

**“Selection Criteria For Patients With Coronary Artery Disease
For Treatment With rFGF-2”**

5

The following selection criteria were applied to Phase I patients with coronary artery disease, whose activities were limited by coronary ischemia despite optimal medical management, and who were not candidates for approved revascularization therapies:

10

Inclusion criteria: Subject is eligible if:

- Male or female, greater than or equal to 18 years of age
- Diagnosis of coronary artery disease (CAD)
- 15 • Suboptimal candidates for approved revascularization procedures, *e.g.*, angioplasty, stents, coronary artery bypass graft (CABG) (or refuses those interventions)
 - Able to exercise at least three minutes using a modified Bruce protocol and limited by coronary ischemia
 - 20 • Inducible and reversible defect of at least 20% myocardium on pharmacologically stressed thallium sestamibi scan
 - CBC, platelets, serum chemistry within clinically acceptable range for required cardiac catheterization
 - Normal INR, or if anticoagulated with Coumadin, INR <2.0
 - 25 • Willing and able to give written informed consent to participate in this study, including all required study procedures and follow-up visits

25

Exclusion criteria: Subject is not eligible if:

30

- Malignancy: any history of malignancy within past ten years, with the exception of curatively treated basal cell carcinoma.
- Ocular conditions: proliferative retinopathy, severe non-proliferative retinopathy, retinal vein occlusion, Eales' disease, or macular

edema or funduscopy by ophthalmologist: history of intraocular surgery within 2 months

- Renal function: creatinine clearance below normal range adjusted for age; protein >250 mg or microalbumin >30 mg/24 h urine
- 5
- Class IV heart failure (New York Heart Association)
 - Ejection fraction < 20% by echocardiogram, thallium scan, MRI or gated pooled blood scan (MUGA)
 - Hemodynamically relevant arrhythmias (*e.g.*, ventricular fibrillation, sustained ventricular tachycardia)
- 10
- Severe valvular stenosis (aortic area <1.0 cm², mitral area <1.2 cm²), or severe valvular insufficiency
 - Marked increase in angina or unstable angina within three weeks
 - History of myocardial infarction (MI) within three months
 - History of transient ischemic attack (TIA) or stroke within 2
- 15 months
- History of CABG, angioplasty or stent within 2 months
 - History of treatment with transmyocardial laser revascularization, rFGF-2, or vascular endothelial growth factor (VEGF) within 2 months
 - Females of child-bearing potential or nursing mothers
- 20
- Any pathological fibrosis, *e.g.*, pulmonary fibrosis, scleroderma
 - Known vascular malformation, *e.g.*, AV malformation, hemangiomas
 - Coexistence of any disease which might interfere with assessment of symptoms of CAD, *e.g.*, pericarditis, costochondritis, esophagitis, systemic
- 25
- vasculitis, sickle cell disease
 - Coexistence of any disease which limits performance of modified Bruce protocol exercise stress test, *e.g.*, paralysis or amputation of a lower extremity, severe arthritis or lower extremities, severe chronic obstructive pulmonary disease (COPD)

- Participation in clinical trials of investigational agents, devices or procedures within thirty days (or scheduled within 60 days of study drug)
 - Known hypersensitivity to rFGF-2 or related compounds
 - Any condition which makes the subject unsuitable for participation
- 5 in this study in the opinion of the investigator, *e.g.*, psychosis, severe mental retardation, inability to communicate with study personnel, drug or alcohol abuse.

EXAMPLE 3

“Phase I Clinical Study on Recombinant FGF-2 (SEQ ID NO: 2) Administered to Humans”

10 The Phase I CAD trial of this example is an open label, dose
15 escalation study of recombinant fibroblast growth factor-2 (rFGF-2) for safety, tolerability and pharmacokinetics. The study was conducted at 2 sites: Beth Israel Deaconess Hospital (Harvard) in Boston, MA and Emory University Hospital in Atlanta, GA. Enrollment is complete. Subjects were treated with a single infusion of rFGF-2 on Day 1 and followed for 360 days; follow-up is not
20 yet complete in some subjects.

The subject population consists of patients with advanced CAD who are exercise limited by coronary ischemia and are considered suboptimal candidates for (or do not want to undergo) one of the established revascularization procedures (*e.g.*, CABG, angioplasty -- with or without stent).
25 The major exclusion criteria were history or suspicion of malignancy, uncompensated heart failure or left ventricular ejection fraction <20%, renal insufficiency or proteinuria, and various ocular conditions (*e.g.*, proliferative diabetic retinopathy, severe non-proliferative retinopathy).

Sixty-six subjects have received rFGF-2 of SEQ ID NO: 2 in this
30 trial: 52 received the rFGF-2 as an IC infusion and fourteen received it as an IV infusion. Each subject was observed in the hospital for at least twenty-four

hours, and then followed as an outpatient for 360 days with follow-up visits (Days 15, 29, 57, 180 and 360). At least four subjects were studied at each dose; if no subject experienced dose-limiting toxicity as defined by the protocol within 2 days, the dose was escalated. The drug was administered as a single 20
5 minute infusion divided between 2 major sources of coronary blood supply (IC), using standard techniques for positioning a catheter into the patient's coronary artery (such as already employed in angioplasty) or in a peripheral vein (IV). The doses in $\mu\text{g/kg}$ of rFGF-2 administered IC (and the number of patients) were 0.33 (n=4), 0.65 (n=4), 2.0 (n=8), 6.0 (n=4), 12.0 (n=4), 24 (n=8), 36
10 (n=10) and 48 (n=10) of rFGF-2 of SEQ ID NO: 2. The doses in $\mu\text{g/kg}$ of rFGF-2 administered IV (and the number of patients) were 18.0 (n=4) and 36.0 (n=10) or rFGF-2 of SEQ ID NO: 2.

Angina frequency and quality of life was assessed by the Seattle Angina Questionnaire (SAQ) at a baseline (before rFGF-2 administration) and at 2
15 months and 6 months after rFGF-2 administration. Exercise tolerance time (ETT) was assessed by the treadmill test at 1, 2, and 6 months. Rest/exercise nuclear perfusion and gated sestamibi-determined rest ejection fraction (EF), and resting magnetic resonance imaging (MRI) were assessed at baseline, and at 1 month, 2 months and 6 months post rFGF-2 administration. MRI measurements which
20 were thought to objectively measure changes in % in cardiac function and perfusion included: (1) ejection fraction; (2) myocardial infarct extent (%); (3) normal wall thickening (4) normal motion (%); (5) target wall thickening (%); (6) target wall motion (%); (7) target wall area collateral extent (%); and (8) target area delayed arrival extent (%).

25 If one of four subjects experienced dose-limiting toxicity (as defined by the protocol), four additional subjects were studied at that dose; if none experienced toxicity, the dose was escalated and another group was studied. The maximally tolerated dose (MTD) was defined as the IC dose which was tolerated by 9/10 subjects, *i.e.*, 36 $\mu\text{g/kg}$ IC.

Careful fluid management pre-infusion was prescribed using a Swan-Ganz catheter and vital signs were monitored frequently during dosing. Heparin was administered IV prior to the infusion of rFGF-2 in all groups. The EDTA concentration was less than 100 µg/ml in the unit dose composition.

- 5 Volume of study drug administered varied with dose and subject's weight, and ranged from 10 ml at lower doses to 40 ml at higher doses.

Preliminary Results

- 10 The results presented here are unaudited and are based on a third interim analysis for 66 subjects with 2 months follow - up for all groups (1-10) and the serious adverse events (SAE) report of 29 July 1999 from Chiron Drug Safety. Data collection for the last visit (Day 360) and final analysis is in progress.

- 15 The starting dose of 0.33 µg/kg IC was escalated over eight sequential groups to 48 µg/kg IC, at which dose 2 of ten subjects experienced dose-limiting toxicity (hypotension) as defined by the protocol. Hypotension was manageable with fluids alone in all subjects (no vasopressors or assistive devices). At 36 µg/kg IC, only 1 of 10 subjects had dose-limiting toxicity which
20 defined this dose as the maximally tolerated dose (MTD). Two additional groups were studied by IV infusion; four subjects of half the MTD (18 µg/kg) and ten subjects at the MTD (36 µg/kg).

- Hypotension was dose-limiting in humans, as predicted by the animal model in Yorkshire pigs. However, 36.0 µg/kg rFGF-2 IC was tolerated
25 in humans; whereas in pigs, 20.0 µg/kg rFGF-2 IC caused profound hypotension in one of 2 animals. Better tolerability in humans was attributed to aggressive fluid management and absence of general anesthesia.

- As of 29 July 1999, thirty-three serious adverse events (SAEs) have occurred in 24/66 subjects, but were not dose-related. Fifteen (15) SAEs
30 were considered at least possibly related to rFGF-2; whenever there was a

difference between relatedness assigned by the investigator and the medical monitor, the more conservative relationship was assigned. SAE's were multiple in five subjects: 01103 (0.33 µg/kg IC), 01106 (0.65 µg/kg IC), 01113 (2.0µg/kg IC), 01137 (36.0 µg/kg IV), 02101 (0.65 µg/kg IC).

5 The most frequent treatment-emergent adverse events (AEs) on Day 1 were transient systolic hypotension and transient bradycardia. The hypotension was dose-dependent and occurred more frequently at doses greater than or equal to (\geq) 24 µg/kg IC; bradycardia was not dose-dependent. Other adverse events (AEs) which were deemed at least possibly related and appeared
10 dose-related occurred within the first several days or week post infusion and included chest pain, shortness of breath, insomnia, anxiety, and nausea. These events were mild to moderate in severity, and most did not require specific intervention.

 When administered IC, the drug was administered over
15 approximately 20 minutes as a single infusion divided between 2 major sources of coronary blood supply (IC), using standard techniques for positioning a catheter into the patient's coronary artery (such as already employed in angioplasty). When administered IV, the drug was administered as an infusion over 20 minutes into a peripheral vein.

20 The preliminary safety results indicate that serious events were not dose related. Thus far, of the eight IC dosage groups, there were three deaths in the lower dosage groups, *i.e.*, at 0.65 µg/kg (Day 23), at 2.0 µg/kg (Day 57) and at 6.0 µg/kg (Day 63), and one death in the highest dose group, *i.e.*, at 48.0 µg/kg (approximately 4 months post-dosing). Three of the deaths were cardiac;
25 one death was due to a large B cell lymphoma that was diagnosed three weeks after dosing in the patient in group 4 (6.0 µg/kg) who patient died at 2 months post-dosing.

 Acute myocardial infarction (MI) occurred in four patients, *i.e.*, one patient from each of groups 1 (0.33 µg/kg), 3 (2.0 µg/kg), 4 (6.0 µg/kg) and 7

(36.0 µg/kg). Multiple MIs occurred in 2 patients, *i.e.*, one from group 1 (0.33 µg/kg) and one from group 3 (2.0 µg/kg). Emergency revascularization procedures (CABG or angioplasty with or without stent) were performed during follow-up in 4 patients: one each from groups 1 (0.33 µg/kg), 3 (2.0 µg/kg), 4
5 (6.0 µg/kg), and 7 (36.0 µg/kg).

Acute hypotension, seen at higher doses during or just subsequent to infusion, was managed by administration of IV fluids without need for a vasopressor. The maximally tolerated dose (MTD) in humans was defined as 36 µg/kg IC. (In contrast, in pigs, the MTD was 6.5 µg/kg IC.) Doses of rFGF-2
10 up to 48 µg/kg IC were administered in human patients with aggressive fluid management, but were defined by the protocol as “not tolerated” due to acute and/or orthostatic hypotension in 2 out of ten patients. The terminal half-life of the infused rFGF-2 was estimated at 5 to 7 hours.

The human patients in this study that were treated with a single IC
15 or IV infusion of rFGF-2 of SEQ ID NO: 2 exhibited a mean increase in ETT of 1.5 to 2 minutes. *See* Table 1. This is especially significant because an increase in ETT of greater than (>) 30 seconds is considered significant and a benchmark for evaluating alternative therapies, such as angioplasty. The angina frequency and quality of life, as measured by SAQ, showed a significant improvement at 2
20 months in all five subscales for the 66 patients (n=66) tested. *See* Tables 2-6. In Tables 2-6, a mean change of 14 or more was considered “clinically significant.”

When 33 human CAD patients were assessed by resting cardiac magnetic resonance imaging (MRI) at baseline, and at 1, 2, and 6 months after receiving a single unit dose composition of the present invention by IC or IV
25 routes, a highly statistically significant increase was observed in target wall thickening, target wall motion and target area collateral extent; a highly statistically significant decrease was observed in target area delayed arrival extent; and no statistically significant changes were observed in normal wall motion, normal wall thickening or myocardial infarct extent.

In addition to the above criterion (*i.e.*, ETT SAQ, MRI), a treatment is considered very successful if the angiogenic effects last at least 2 months. In the present Phase I study, the unexpectedly superior angiogenic effects were observed to last up to 6 months in some patients in all dosage groups. Based upon the results already obtained, it is expected that the angiogenic effects may last twelve months or more but do last at least 2 months in the patients, at which time the procedure could be repeated, if necessary.

EXAMPLE 4

“Proposed Phase II Clinical Study On rFGF-2 (SEQ ID NO: 2) Administered to Humans to Treat Coronary Artery Disease”

The Phase II clinical trial of rFGF-2 for treating human patients for coronary artery disease is performed as a double blind/placebo controlled study having four arms: placebo, 0.3 µg/kg, 3.0 µg/kg, and 30 µg/kg administered once IC. This study is ongoing and results are not yet available.

EXAMPLE 5

“Unit Dose and Pharmaceutical Composition of rFGF-2 for the Phase II Human Clinical Trial”

The rFGF-2 of SEQ ID NO: 2 was formulated as a unit dose pharmaceutical composition for administration to humans in the Phase II clinical trial referenced herein. The various formulations are described below.

The rFGF-2 Unit Dose was prepared as a liquid in 5 cc type I glass vials with a laminated gray butyl rubber stopper and red flip-off overseal. The rFGF-2 formulation contains 0.3 mg/ml rFGF-2 of SEQ ID NO: 2 in 10 mM sodium citrate, 10 mM monothioglycerol, 0.3 mM disodium dihydrate EDTA (molecular weight 372.2), 135 mM sodium chloride, pH 5.0. Each vial contained 3.7 ml of rFGF-2 drug product solution (1.11 mg rFGF-2 per vial). The resulting unit dose in liquid form is stored at less than -60° C. The above described unit dose

is diluted with the “rFGF-2 placebo.” Depending on the size of the patient, the contents of several of the vials may be combined to produce a unit dose of 36 µg/kg for the Phase II study.

- 5 **The rFGF-2 placebo** is supplied as a clear colorless liquid in 5 cc type I glass vials with a laminated gray butyl rubber stopper and red flip-off overseal. The rFGF-2 placebo is indistinguishable in appearance from the drug product and has the following formulation: 10 mM sodium citrate, 10 mM monothioglycerol, 0.3 mM disodium dihydrate EDTA (molecular weight 372.2), 135 mM sodium chloride, pH 5.0. Each vial contains 5.2 ml of rFGF-2 placebo solution. Unlike
10 the unit dose, the rFGF-2 placebo is stored at 2° to 8°C.

- 15 **The rFGF-2 Pharmaceutical Composition** that is infused is prepared by diluting the rFGF-2 unit dose with the rFGF diluent such that the infusion volume is 20 ml for Phase II.

Example 6

Primary Objective

- 20 To compare the effect of a single, IC infusion of rFGF-2 versus placebo on exercise capacity, as measured by the change in exercise tolerance test (ETT) time from baseline (during the screening period) to 90 days.

Secondary Objectives

- To evaluate the safety of rFGF-2, as measured by AEs and changes in laboratory parameters.
- 25 • To confirm the pharmacokinetics (PK) of rFGF-2 from the Phase I study.
- To evaluate the effect of rFGF-2 on:
 - Change in ETT time from baseline to 180 days.

- Changes from baseline in quality of life (QoL): perceived angina as measured by the Seattle Angina Questionnaire (SAQ) and general well-being, as measured by the Short Form-36 (SF-36) at 90 and 180 days.
- Changes from baseline in the size of the ischemic area at rest and with pharmacologic stress with dipyridamole by thallium/sestamibi scans at 90 and 180 days.
- In a subset of subjects, to evaluate the effect of rFGF-2 on EF, target wall motion, and wall thickness and perfusion by magnetic resonance imaging (MRI) scan.

10 INVESTIGATIONAL PLAN

OVERALL STUDY DESIGN AND PLAN

- This clinical trial was designed as a phase 2, multicenter, double-blind, placebo-controlled study of rFGF-2 in 300 subjects with CAD. Subjects who met all eligibility criteria were randomly assigned to receive placebo or one of three doses of rFGF-2 (approximately 75 subjects per arm). Doses were 0 (placebo), 0.3, 3.0, and 30.0 µg/kg based on actual body weight. Study drug was to be administered as a single IC infusion over 20 minutes during cardiac catheterization. Subjects were monitored in the hospital for at least 6 hours postdose at the site, and then followed at specified intervals over 180 days. Long-term safety was assessed in a separate extension protocol using telephone contacts and questionnaires for an additional 6 months.

Inclusion Criteria

The following inclusion criteria were required for study entry:

- Men or women ≥ 18 years of age.
- Diagnosis of CAD, as defined by $> 60\%$ stenosis of a major coronary artery by coronary angiography.

- Ejection fraction $\geq 30\%$ by an accepted imaging technology (ventriculogram, MRI, first pass on single photon emission computed tomography [SPECT]) scan, echocardiogram [ECHO] or multigated [MUGA] nuclear assessment).
 - Symptoms of angina or angina equivalent.
- 5
- Relegated to non-invasive therapy for CAD under the care of a physician.
 - Not a candidate for standard surgical or catheter-based revascularization procedures as assessed by a physician.
 - No evidence of malignancy on mammography within 12 months (females).
 - Pelvic exam with cervical smear within 1 year of screening for all females.
- 10
- Sigmoidoscopy within 5 years for all subjects ≥ 50 years.
 - Annual rectal exam with stool guaiac for subjects ≥ 40 years.
 - Able to exercise on baseline treadmill exercise testing for at least 3 minutes but not longer than 13 minutes using a modified Bruce protocol, and limited by signs or symptoms of ischemia. Subjects had to complain of some degree of
- 15
- angina or anginal equivalent or have ≥ 1 mm ST segment depression during the course of the ETTs. Duplicate baseline tests were performed at least 24 hours but not greater than 2 weeks apart, and the difference between the two tests had to be $\leq 20\%$ of the average of the two tests.
- 20
- Inducible and reversible ischemic defect of moderate or greater size involving at least half the area of one major myocardial territory (anterior, inferior, lateral, or septal) on a rest/stressed thallium/sestamibi scan, or multiple inducible and reversible ischemic defects in multiple myocardial territories, the sum total of which involves the equivalent of at least half the area of one major myocardial territory on a rest/stressed thallium/sestamibi scan within 30 days
- 25
- prior to randomization.
 - Complete blood count (CBC), platelets, serum chemistry, prothrombin time, and urinalysis within clinically acceptable ranges for cardiac catheterization within 30 days prior to randomization.

- Protein on a urine sample negative or trace within 30 days prior to randomization.
- Serum creatinine ≤ 2.0 mg/dL within 30 days prior to randomization.
- Willing and able to give written informed consent to participate in this study, including all required study procedures and follow-up visits.
- Prostate specific antigen (PSA) within 30 days prior to randomization (males) conforming to the following age-specific ranges:

	<u>Age</u>	<u>PSA (ng/mL)</u>
	<40 years	< 2.0
10	40-49 years	< 2.7
	50-59 years	< 3.7
	60-69 years	< 5.1
	70-79 years	< 7.0
	≥ 80 years	< 7.2

- No evidence of malignancy on chest x-ray within 30 days prior to randomization
- No evidence or suspicion of malignancy as assessed by complete history and physical exam (performed according to American Cancer Society guidelines) within 30 days of randomization.
- Confirmation on Study Day 1 by cardiac catheterization that the subject's cardiovascular anatomy is not suitable for treatment with an invasive procedure according to the opinion of the principal investigator or another study physician at the site.

25

Exclusion Criteria

- Malignancy: History or suspicion of malignancy within the past 10 years, with the exception of curatively treated basal cell carcinoma, squamous cell carcinoma of the skin in sun-exposed areas, or carcinoma of the cervix.
- Renal conditions:

30

- Renal insufficiency, as defined by serum creatinine > 2.0 mg/dL.
- Proteinuria, as defined by 1+ or greater protein on dipstick.
- Ocular conditions:
 - Proliferative retinopathy, moderate or severe nonproliferative retinopathy.
- 5
 - Retinal vein occlusion.
 - Age-related maculopathy with choroidal neovascularization.
 - Macular edema on funduscopy by an ophthalmologist.
 - Intraocular surgery within 4 months.
- Cardiovascular conditions:
 - 10
 - Severe aortic stenosis, ie, area < 1.0 cm².
 - Unstable angina within 3 weeks (Braunwald, 1984).
 - CABG, angioplasty, transient ischemic attack, or stroke within 4 months.
 - Myocardial infarction within 3 months.
 - Treatment with transmyocardial laser revascularization within 1 year of
 - 15 screening.
 - Current treatment with external counterpulsation.
 - Past participation in any therapeutic angiogenesis trial with any investigational agent unless it could be demonstrated that the subject definitely received placebo.
 - 20
 - Diagnosis of primary pulmonary hypertension or restrictive or obstructive cardiomyopathy.
- General medical conditions:
 - Pregnancy or nursing mothers.
 - Any pathological fibrosis, eg, pulmonary fibrosis, scleroderma.
 - 25
 - Known vascular malformation, eg, arteriovenous malformation, hemangiomas >3 mm.
 - Participation in clinical trials of other investigational agents, IC devices or procedures, for which follow-up visits had not been completed.
 - History of organ transplantation.

- Any combined condition which made the subject unsuitable for participation in this study in the opinion of the investigator, eg, concurrent medical illness which limits life expectancy to < 12 months, psychosis, severe mental retardation, inability to communicate with study personnel, drug or alcohol abuse.
- 5

Removal of Subjects From Therapy or Assessment

- A subject could withdraw consent to participate in the study at any time without prejudice. Additionally, the investigator could withdraw a subject if, according to clinical judgment, it was in the best interest of the subject or if the subject could not comply with the protocol.
- 10

If a subject who met all inclusion and exclusion requirements and received study drug on Day 1 subsequently withdrew from the study, the tests and evaluations listed for the day 180 termination visit were carried out whenever possible.

- 15 If a subject developed an AE that the investigator believed was severe enough to interrupt study drug administration for > 10 minutes, dosing was discontinued and the subject was not to be allowed to receive further study drug treatment. The subject was to receive medical treatment as determined by the investigator, remain under observation until the AE resolved or became stable, and
- 20 was to be followed through day 180.

Treatments Administered

Subjects were randomly assigned to receive placebo or 0.3, 3.0, or 30.0 $\mu\text{g/kg}$ rFGF-2. The study drug was to be administered as an infusion of 20 mL over 20 minutes divided between two patent coronary vessels or grafts, using a calibrated, precision, infusion pump.

Identity of Investigational Product

The rFGF-2 used in this study was a 146 amino acid, non-glycosylated, monomeric, 16.5 kDa protein that was expressed in genetically engineered yeast.

The rFGF-2 drug product was supplied as a clear, colorless liquid in 5-mL, type I, glass vials with a laminated, gray, butyl stopper and red, flip-off overseal. The rFGF-2 formulation contained 0.3 mg/mL rFGF-2 in 10 mM sodium citrate, 10 mM monothioglycerol, 0.3 mM disodium dihydrate EDTA (molecular weight 372.2), 135 mM sodium chloride, pH 5.0. Each vial contained 3.7 mL of rFGF-2 drug-product solution (1.11 mg rFGF-2 per vial). The rFGF-2 drug-product vials were stored at -60°C or less. The rFGF-2 was diluted with placebo according to the subjects' actual body weight.

The placebo was supplied as a clear, colorless liquid in 5-mL, type I, glass vials with a laminated, gray, butyl stopper and red, flip-off overseal. The placebo was indistinguishable in appearance from the drug product; it contained 10 mM sodium citrate, 10 mM monothioglycerol, 0.3 mM disodium dihydrate EDTA (molecular weight 372.2), 135 mM sodium chloride, pH 5.0. Each vial contained 5.2 mL of placebo solution. The placebo vials were stored at 2 to 8°C .

Selection of Doses in the Study

The doses selected were those that bracketed a range of doses for which preclinical and early clinical data suggested the best probability of efficacy. The safety of the highest dose (30.0 $\mu\text{g/kg}$) was supported by the phase 1 trial results

reported in Example 3.

Subjects typically received a single IV bolus of heparin between 10 and 20 minutes prior to infusion of study drug in order to minimize the risk of thrombosis related to the duration of time the catheter was in place.

5 EFFICACY AND SAFETY VARIABLES

Efficacy and Safety Measurements Assessed and Flowchart

The primary efficacy variable was the change in exercise capacity as measured by ETT time at Day 90. Secondary efficacy variables included the change in ETT time at Day 180; the change in QoL as measured by the angina frequency score (AFS), treatment satisfaction score (TSS), exertional capacity score (ECS), and disease perception score (DPS) of the SAQ, and the physical and mental components of the SF-36 at Days 90 and 180; the change in ischemic area at rest and with pharmacologic stress by thallium/sestamibi scans at Days 90 and 180; changes in EF, targeted wall thickness and motion, and perfusion by MRI at Days 90 and 180 in a subset of subjects.

Other analysis variables added at the time that the statistical analysis plan was implemented included time to onset of angina, time to onset of 1-mm ST segment depression, and changes from baseline in rate-pressure product at onset of angina, in rate-pressure product at peak exercise, and in rate-pressure product at 1-mm ST segment depression at Days 90 and 180. Changes in the New York Heart Association (NYHA) classification and the Canadian Cardiovascular Class (CCS) were analyzed. The percentage of subjects with ≥ 60 seconds increase in ETT time (“responder”) and of subjects with < 60 seconds increase in ETT time (“non-responder”) was analyzed for each group.

The Canadian Cardiovascular Classification is based on a classification scheme ranging from Class 0 to Class IV. To be classified in: Class 0, a patient

does not experience angina or anginal equivalent symptoms; Class I, with ordinary physical activity symptoms occur only with strenuous, rapid, or prolonged exertion at work or recreation; Class II, a patient experiences slight limitation of ordinary activity due to angina; Class III, a patient experiences marked limitation of activity
5 due to angina; and Class IV, a patient develops angina at rest or with any physical activity.

The SAQ is a validated, disease-specific, self-administered questionnaire with 5 scales : angina stability scale (ASS), angina frequency scale (AFS), exercise capacity scale (ECS), treatment satisfaction scale (TSS), and
10 disease perception scale (DPS) (Spertus JA, et al.). The ASS was not included in the analysis of this trial as it refers to a 4 week interval which was not relevant to the evaluation periods. Each component comprises a scale from 1 to 100; lower scores are associated with worse symptoms; ≥ 8 point changes are considered clinically relevant. The SF-36 is a validated, general, quality-of-life (QoL)
15 instrument with 10 scales and two summary component scales: a physical component summary scale (PCSS) and a mental component summary scale (MCSS).

Nuclear imaging results were analyzed using a validated semiquantitative grading system in a 20-segment left ventricular model (Berman
20 DS, et al.). The perfusion of each segment was graded on a five-point scale: 0 = normal, 1 = slightly reduced, 2 = moderately reduced, 3 = severely reduced, and 4 = absent. A segment was determined to have a reversible defect if the assigned abnormal regional grade at stress decreased or normalized on the rest images (reversibility score: stress score – rest score >1), and to have a fixed defect if the
25 assigned regional grade at stress was abnormal and remained the same on rest imaging. The fixed defects were subgrouped on the basis of the severity of the graded scores, i.e., mild-to-moderate (scores of 1, 2, and 3) and severe (score of 4). The global extent of perfusion abnormality and ischemia was assessed by

summing the individual scores from the 20 segments, and expressed as the average stress and average reversibility scores, respectively.

Safety was monitored by evaluating AEs, laboratory data, and the results of physical examinations. A Data and Safety Monitoring Board (DSMB) reviewed SAEs and results of laboratory tests approximately every 6 weeks. Acute cardiac events were adjudicated by a Clinical Events Committee (CEC). Core laboratories reviewed the results of angiograms, ECGs, ophthalmologic evaluations, and nuclear scans.

DEMOGRAPHIC AND OTHER SUBJECT CHARACTERISTICS

Table 7 summarizes demographic features of the subjects enrolled in the trial. These features were similar among the four treatment groups (Table 7).

Table 7 Summary of Demography

Treatment Group	Placebo	0.3 µg/kg	3.0 µg/kg	30 µg/kg
Age (mean years)	63.9	63.0	62.9	61.8
Percent male sex	86%	84%	80%	86%
Percent Caucasian race	97%	90%	94%	93%
Weight (mean kg)	88.24	87.25	87.23	87.50
Height (mean cm)	170.94	172.02	172.45	171.99
Percent tobacco use	8%	11%	10%	10%

Analysis

Exercise Tolerance Testing

Figure 9 shows the ETT time at baseline, at Day 90, and the primary efficacy variable, change in ETT time from baseline to Day 90. In this and the tabulations of all efficacy variables, subjects who had undergone a revascularization and subjects with no assessment were excluded from the analysis unless otherwise specified. The mean increase in exercise time was 44.1 seconds for the placebo group, 48.5 seconds in the low-dose group, 65.0 seconds in the middle-dose group, and 49.1 seconds in the high-dose group.

Figure 9 also shows ETT time at Day 180 and the change in ETT time from baseline to Day 180. The mean increase in exercise time was 54.8 seconds in the placebo group, 75.3 seconds in the low-dose group, 76.3 seconds in the middle-dose group, and 51.3 seconds in the high-dose group. Trends in improvement in ETT were observed at Day 180 in subjects with a baseline CCS of 3 or 4 and/or SAQ angina frequency of less than or equal to 40.

Seattle Angina Questionnaire

Figure 10 shows the analyses of the SAQ. A change of ≥ 8 points is considered clinically meaningful and higher SAQ scores are associated with better clinical status. For AFS change scores at Day 90, the P values were .035 based upon the overall test and .007 based upon the test of placebo versus all FGF. The mean change from baseline was 8.1 for the placebo group, 16.0 for the low-dose group, 20.8 for the middle-dose group, and 16.7 for the high-dose group (pairwise P values = .080, .004, .054, respectively). For AFS change scores at Day 180, the mean change was 15.3 for the placebo group, 18.7 for the low-dose group, 22.6 for the middle-dose group, and 20.2 for the high-dose group (pairwise P values = .44, .089, .25, respectively). The remaining SAQ scales, Exertional Capacity, Treatment Satisfaction, and Disease Perception, are shown for the FGF treatment groups in Figure 11.

Short Form 36

Figure 12 shows the analysis for the SF-36 Physical Component Summary Scale (PCSS) and Mental Component Summary Scale (MCSS). For the PCSS at Day 90, the P value based on the test of placebo versus all FGF was .033. The mean change from baseline was 3.4 in the placebo group, 6.0 in the low-dose group, 5.9 in the middle-dose group, and 5.9 in the high-dose group (pairwise P values = .072, .091, .095, respectively). For PCSS at Day 180, the difference in the

mean change from baseline was less than 2.2 between the placebo and FGF groups. Treatment effect was not detectable in the MCSS (mental component) of the SF-36.

Nuclear Imaging

Consistent differences among the placebo and FGF groups were not detectable.

5 Examination of Subgroups

Exercise Tolerance Testing

Figure 13 (left panel) shows ETT time for subjects with baseline CCS of 3 or 4. The mean change in exercise time from baseline to Day 90 for subjects with a baseline CCS of 3 or 4 was 36.1 seconds for the placebo group, 47.6 seconds for the low-dose group, 57.1 seconds for the middle-dose group, and 44.5 seconds in the high-dose group (pairwise *P* values = .59, .31, .69, respectively). The mean change in exercise time from baseline to Day 180 was 33.1 seconds for the placebo group, 70.7 seconds for the low-dose group, 75.7 seconds for the middle-dose group, and 41.5 seconds for the high-dose group (pairwise *P* values = .15, .086, .74, respectively).

Figure 13 (right panel) show SAQ scores for subjects with baseline CCS of 3 or 4. The mean change in AFS from baseline to Day 90 for subjects with a baseline CCS of 3 or 4 was 5.7 for the placebo group, 17.0 for the low-dose group, 19.5 for the middle-dose group, and 20.5 for the high-dose group (pairwise *P* values = .058, .015, .010, respectively). At Day 180, the mean change in AFS was 11.7 for the placebo group, 17.2 for the low-dose group, 22.3 for the middle-dose group, and 21.8 for the high-dose group (pairwise *P* values = .34, .059, .076, respectively).

Figure 14 (left panel) shows ETT time for subjects with baseline Angina Frequency Score (AFS) of ≤ 40 , as measured using the Seattle Angina

- Questionnaire (SAQ). The mean change in exercise time from baseline to Day 90 for subjects with a baseline AFS ≤ 40 (lower scores indicating a higher frequency of angina) was 50.3 seconds for the placebo group, 83.6 seconds for the low-dose group, 72.1 seconds for the middle-dose group, and 49.8 seconds for the high-dose group (pairwise P values = .16, .32, .98, respectively). The mean change in exercise time from baseline to Day 180 was 56.5 seconds for the placebo group, 90.2 seconds for the low-dose group, 84.6 seconds for the middle-dose group, and 55.0 seconds for the high-dose group (pairwise P values = .24, .29, .96, respectively).
- 10 These data suggest a higher mean change in ETT time for subjects with a higher frequency of angina at baseline and who received the low or middle doses of FGF compared to those who received placebo.

- Figure 14 (right panel) shows AFS for subjects with baseline AFS of ≤ 40 . The mean change in AFS from baseline to Day 90 for subjects with a baseline AFS ≤ 40 (lower scores indicating a higher frequency of angina) was 12.2 for the placebo group, 27.7 for the low-dose group, 29.7 for the middle-dose group, and 22.0 for the high-dose group (pairwise P values = .017, .004, .10, respectively). At Day 180, the mean change in AFS was 16.3 for the placebo group, 27.4 for the low-dose group, 30.7 for the middle-dose group, and 24.6 for the high-dose group (pairwise P values = .089, .019, .18, respectively). The magnitude of the difference in AFS between the low and middle dose groups and placebo in subjects with lower AFS at baseline was considered clinically relevant.
- 15
- 20

Functional Analysis

25 Anatomy

At 90 days there was a significant reduction in Left Ventricular End Diastolic Diameter (LVEDD) in the 3.0 ug/kg dosage group and in the 30 ug/kg

dosage group ($p=0.037$ and 0.032 , respectively). There was a trend to improvement in the 0.3 ug/kg dosage group ($p=0.13$). In controls, LVEDD slightly increased. This is consistent with relief of ischemia and an improvement in function in FGF treated patients (3.0 ug/kg and 30 ug/kg dose groups). At 90 days there is a significant reduction in Left Ventricular End Systolic Diameter (LVESD) in 0.3 ug/kg dose group ($p=0.042$) and the 30 ug/kg dose group ($p=0.024$) and a trend ($p=0.17$) in the 3.0 ug/kg group. This is also consistent with a relief of ischemia and an improvement in function in FGF-treated patients. Controls showed an increase in LVESD (consistent with an increase in LVEDD). The effect is not evident at 6 months.

Left Ventricular (LV) mass was unchanged, as would be expected (Because there was no effect on blood pressure, no effect on LV mass was expected.).

Wall motion

Normal wall motion is in the expected range (30-40%) and was unchanged. Normal wall thickening is in the expected range (45-50%) and is unchanged. Target wall, i.e., the wall of the myocardium that is ischemic, motion at rest was the same as normal.

Myocardial perfusion

Delayed arrival zone (DAZ) was similar across all groups at baseline (mean~15%). That correlated with the size of exercise-induced nuclear perfusion defect (~18%). The size of DAZ declines in all groups. The highest decline was the 3.0 ug/kg group.

Summary and Conclusions:

The data from the Phase II trial showed an overall patterns that suggests that subjects improve most with the low (0.3 ug/kg) and mid-doses (3.0 ug/kg) of FGF-2.

Overall, more symptomatic patients showed the most improvement. For example, subjects with a baseline of CCS of 3 or 4 and/or SAQ Angina Frequency of ≤ 40 showed a trend towards improvement at ETT at Day 180.

For all subject groups, reduction of angina frequency ($p=0.035$) was observed at Day 90. The subjects with a baseline of CCS of 3 or 4 and/or SAQ Angina Frequency of ≤ 40 showed the most improvement at both Day 90 and Day 180. Further, subjects with a baseline of CCS of 3 or 4 showed improvement in the SF-36 Physical Component at both Day 90 and Day 180.

Further, MRI data provided evidence of improved LV function in FGF groups as evidenced by reduced LVEDD and LVESD. Perfusion analysis showed a trend towards improvement in FGF groups.

For the secondary efficacy variables of the SAQ, changes ≥ 8 are considered clinically relevant; a statistically significant difference was seen in the overall analysis of the angina frequency score (AFS). Differences that were clinically relevant and statistically significant were seen between the placebo and the middle-dose groups in AFS at Day 90.

For the secondary efficacy variables of the SF-36, a significant difference between the placebo and all FGF-treated groups was seen in the physical component summary scale (PCSS) at Day 90 (0.033).

Additionally, in a subset of patients with congestive heart failure, improvement was seen with any dose of FGF. Thus, FGF may be more helpful in patients with congestive heart failure. In the Phase II study, at day 90, patients receiving placebo showed a change of 23.2 seconds (least square mean) from baseline for total exercise time as compared with patients receiving low doses and mid-range doses who showed a change of 64.6 and 81.3, respectively. At day 180, the placebo showed a change in total exercise time of 27.3 seconds as compared to

75.4 seconds and 72.8 seconds for the low dose and mid-dose groups. The results establish a statistical trend in mid and low dosage groups at day 90.